

5 The EGR1 gene expression is activated during this treatment. Thus, by stably
transfecting PC12 cells with a construct containing an EGR promoter linked to SEAP
10 reporter, activation of PC12 cells can be assessed.

The EGR/SEAP reporter construct can be assembled by the following
5 protocol. The EGR-1 promoter sequence (-633 to +1)(Sakamoto K et al., Oncogene
6:867-871 (1991)) can be PCR amplified from human genomic DNA using the
15 following primers:

5' GCGCTCGAGGGATGACAGCGATAGAACCCCGG -3' (SEQ ID NO:6)

5' GCGAAGCTTCGCGACTCCCCGGATCCGCCTC-3' (SEQ ID NO:7)

20 Using the GAS:SEAP/Neo vector produced in Example 12, EGR1 amplified
product can then be inserted into this vector. Linearize the GAS:SEAP/Neo vector
using restriction enzymes XhoI/HindIII, removing the GAS/SV40 stuffer. Restrict the
EGR1 amplified product with these same enzymes. Ligate the vector and the EGR1
25 promoter.

15 To prepare 96 well-plates for cell culture, two mls of a coating solution (1:30
dilution of collagen type I (Upstate Biotech Inc. Cat#08-115) in 30% ethanol (filter
sterilized)) is added per one 10 cm plate or 50 ml per well of the 96-well plate, and
30 allowed to air dry for 2 hr.

PC12 cells are routinely grown in RPMI-1640 medium (Bio Whittaker)
20 containing 10% horse serum (JRH BIOSCIENCES, Cat. # 12449-78P), 5% heat-
inactivated fetal bovine serum (FBS) supplemented with 100 units/ml penicillin and
35 100 ug/ml streptomycin on a precoated 10 cm tissue culture dish. One to four split is
done every three to four days. Cells are removed from the plates by scraping and
resuspended with pipetting up and down for more than 15 times.

40 25 Transfect the EGR/SEAP/Neo construct into PC12 using the Lipofectamine
protocol described in Example 11. EGR-SEAP/PC12 stable cells are obtained by
growing the cells in 300 ug/ml G418. The G418-free medium is used for routine
45 growth but every one to two months, the cells should be re-grown in 300 ug/ml G418
for couple of passages.

30 To assay for neuronal activity, a 10 cm plate with cells around 70 to 80%
confluent is screened by removing the old medium. Wash the cells once with PBS
50

(Phosphate buffered saline). Then starve the cells in low serum medium (RPMI-1640 containing 1% horse serum and 0.5% FBS with antibiotics) overnight.

The next morning, remove the medium and wash the cells with PBS. Scrape off the cells from the plate, suspend the cells well in 2 ml low serum medium. Count the cell number and add more low serum medium to reach final cell density as 5×10^5 cells/ml.

Add 200 μ l of the cell suspension to each well of 96-well plate (equivalent to 1×10^5 cells/well). Add 50 μ l supernatant produced by Example 11, 37°C for 48 to 72 hr. As a positive control, a growth factor known to activate PC12 cells through EGR can be used, such as 50 ng/ μ l of Neuronal Growth Factor (NGF). Over fifty-fold induction of SEAP is typically seen in the positive control wells. SEAP assay the supernatant according to Example 17.

Example 16: High-Throughput Screening Assay for T-cell Activity

NF- κ B (Nuclear Factor κ B) is a transcription factor activated by a wide variety of agents including the inflammatory cytokines IL-1 and TNF, CD30 and CD40, lymphotoxin-alpha and lymphotoxin-beta, by exposure to LPS or thrombin, and by expression of certain viral gene products. As a transcription factor, NF- κ B regulates the expression of genes involved in immune cell activation, control of apoptosis (NF- κ B appears to shield cells from apoptosis), B and T-cell development, anti-viral and antimicrobial responses, and multiple stress responses.

In non-stimulated conditions, NF- κ B is retained in the cytoplasm with I- κ B (Inhibitor κ B). However, upon stimulation, I- κ B is phosphorylated and degraded, causing NF- κ B to shuttle to the nucleus, thereby activating transcription of target genes. Target genes activated by NF- κ B include IL-2, IL-6, GM-CSF, ICAM-1 and class I MHC.

Due to its central role and ability to respond to a range of stimuli, reporter constructs utilizing the NF- κ B promoter element are used to screen the supernatants produced in Example 11. Activators or inhibitors of NF- κ B would be useful in treating diseases. For example, inhibitors of NF- κ B could be used to treat those

diseases related to the acute or chronic activation of NF- κ B, such as rheumatoid arthritis.

To construct a vector containing the NF- κ B promoter element, a PCR based strategy is employed. The upstream primer contains four tandem copies of the NF- κ B binding site (GGGGACTTTCCC) (SEQ ID NO:8), 18 bp of sequence complementary to the 5' end of the SV40 early promoter sequence, and is flanked with an XhoI site:
5':GCGGCCTCGAGGGGACTTTCCCGGGGACTTTCCGGGGACTTTCCGGGGACTTTCCGGGGACTTTCCATCCTGCCATCTCAATTAG:3' (SEQ ID NO:9)

The downstream primer is complementary to the 3' end of the SV40 promoter and is flanked with a Hind III site:
5':GCGGCAAGCTTTTGGCAAAGCCTAGGC:3' (SEQ ID NO:4)

PCR amplification is performed using the SV40 promoter template present in the pB-gal:promoter plasmid obtained from Clontech. The resulting PCR fragment is digested with XhoI and Hind III and subcloned into BLSK2-. (Stratagene)
Sequencing with the T7 and T3 primers confirms the insert contains the following sequence:

5':CTCGAGGGGACTTTCCCGGGGACTTTCCGGGGACTTTCCGGGGACTTTCCATCTGCCATCTCAATTAGTCAGCAACCATAGTCCCGCCCCTAACTCCGCCCATCCCGCCCCTAACTCCGCCCAGTTCGCCCCATTCTCGCCCCCATGGCTGACTAATTATTTTATTTATGTCAGAGGCCGAGGCCGCTCGGCCTCTGAGCTATTCCAGAAGTAGTGAGGAGGCTTTTTTGGAGGCCTAGGCTTTTGCAAAAAA GCTT:3' (SEQ ID NO:10)

Next, replace the SV40 minimal promoter element present in the pSEAP2-promoter plasmid (Clontech) with this NF- κ B/SV40 fragment using XhoI and HindIII. However, this vector does not contain a neomycin resistance gene, and therefore, is not preferred for mammalian expression systems.

In order to generate stable mammalian cell lines, the NF- κ B/SV40/SEAP cassette is removed from the above NF- κ B/SEAP vector using restriction enzymes SalI and NotI, and inserted into a vector containing neomycin resistance. Particularly,

the NF- κ B/SV40/SEAP cassette was inserted into pGFP-1 (Clontech), replacing the GFP gene, after restricting pGFP-1 with SalI and NotI.

Once NF- κ B/SV40/SEAP/Neo vector is created, stable Jurkat T-cells are created and maintained according to the protocol described in Example 13. Similarly, the method for assaying supernatants with these stable Jurkat T-cells is also described in Example 13. As a positive control, exogenous TNF alpha (0.1, 1, 10 ng) is added to wells H9, H10, and H11, with a 5-10 fold activation typically observed.

Example 17: Assay for SEAP Activity

As a reporter molecule for the assays described in Examples 13-16, SEAP activity is assayed using the Tropix Phospho-light Kit (Cat. BP-400) according to the following general procedure. The Tropix Phospho-light Kit supplies the Dilution, Assay, and Reaction Buffers used below.

Prime a dispenser with the 2.5x Dilution Buffer and dispense 15 μ l of 2.5x dilution buffer into Optiplates containing 35 μ l of a supernatant. Seal the plates with a plastic sealer and incubate at 65°C for 30 min. Separate the Optiplates to avoid uneven heating.

Cool the samples to room temperature for 15 minutes. Empty the dispenser and prime with the Assay Buffer. Add 50 μ l Assay Buffer and incubate at room temperature 5 min. Empty the dispenser and prime with the Reaction Buffer (see the table below). Add 50 μ l Reaction Buffer and incubate at room temperature for 20 minutes. Since the intensity of the chemiluminescent signal is time dependent, and it takes about 10 minutes to read 5 plates on luminometer, one should treat 5 plates at each time and start the second set 10 minutes later.

Read the relative light unit in the luminometer. Set H12 as blank, and print the results. An increase in chemiluminescence indicates reporter activity.

Reaction Buffer Formulation:

# of plates	Rxn buffer diluent (ml)	CSPD (ml)
10	60	3
11	65	3.25
12	70	3.5

5

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	13	75	3.75
	14	80	4
	15	85	4.25
10	16	90	4.5
	17	95	4.75
	18	100	5
	19	105	5.25
15	20	110	5.5
	21	115	5.75
	22	120	6
	23	125	6.25
20	24	130	6.5
	25	135	6.75
	26	140	7
	27	145	7.25
	28	150	7.5
25	29	155	7.75
	30	160	8
	31	165	8.25
	32	170	8.5
30	33	175	8.75
	34	180	9
	35	185	9.25
	36	190	9.5
35	37	195	9.75
	38	200	10
	39	205	10.25
	40	210	10.5
40	41	215	10.75
	42	220	11
	43	225	11.25
	44	230	11.5
	45	235	11.75
45	46	240	12
	47	245	12.25
	48	250	12.5
	49	255	12.75
50	50	260	13

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Example 18: High-Throughput Screening Assay Identifying Changes in Small Molecule Concentration and Membrane Permeability

Binding of a ligand to a receptor is known to alter intracellular levels of small molecules, such as calcium, potassium, sodium, and pH, as well as alter membrane potential. These alterations can be measured in an assay to identify supernatants which bind to receptors of a particular cell. Although the following protocol describes an assay for calcium, this protocol can easily be modified to detect changes in potassium, sodium, pH, membrane potential, or any other small molecule which is detectable by a fluorescent probe.

The following assay uses Fluorometric Imaging Plate Reader ("FLIPR") to measure changes in fluorescent molecules (Molecular Probes) that bind small molecules. Clearly, any fluorescent molecule detecting a small molecule can be used instead of the calcium fluorescent molecule, fluo-4 (Molecular Probes, Inc.; catalog no. F-14202), used here.

For adherent cells, seed the cells at 10,000 -20,000 cells/well in a Co-star black 96-well plate with clear bottom. The plate is incubated in a CO₂ incubator for 20 hours. The adherent cells are washed two times in Biotek washer with 200 ul of HBSS (Hank's Balanced Salt Solution) leaving 100 ul of buffer after the final wash.

A stock solution of 1 mg/ml fluo-4 is made in 10% pluronic acid DMSO. To load the cells with fluo-4, 50 ul of 12 ug/ml fluo-4 is added to each well. The plate is incubated at 37°C in a CO₂ incubator for 60 min. The plate is washed four times in the Biotek washer with HBSS leaving 100 ul of buffer.

For non-adherent cells, the cells are spun down from culture media. Cells are re-suspended to 2-5x10⁶ cells/ml with HBSS in a 50-ml conical tube. 4 ul of 1 mg/ml fluo-4 solution in 10% pluronic acid DMSO is added to each ml of cell suspension. The tube is then placed in a 37°C water bath for 30-60 min. The cells are washed twice with HBSS, resuspended to 1x10⁶ cells/ml, and dispensed into a microplate, 100 ul/well. The plate is centrifuged at 1000 rpm for 5 min. The plate is then washed once in Denley CellWash with 200 ul, followed by an aspiration step to 100 ul final volume.

For a non-cell based assay, each well contains a fluorescent molecule, such as fluo-4. The supernatant is added to the well, and a change in fluorescence is detected.

To measure the fluorescence of intracellular calcium, the FLIPR is set for the following parameters: (1) System gain is 300-800 mW; (2) Exposure time is 0.4 second; (3) Camera F/stop is F/2; (4) Excitation is 488 nm; (5) Emission is 530 nm; and (6) Sample addition is 50 ul. Increased emission at 530 nm indicates an extracellular signaling event which has resulted in an increase in the intracellular Ca^{++} concentration.

Example 19: High-Throughput Screening Assay Identifying Tyrosine Kinase Activity

The Protein Tyrosine Kinases (PTK) represent a diverse group of transmembrane and cytoplasmic kinases. Within the Receptor Protein Tyrosine Kinase (RPTK) group are receptors for a range of mitogenic and metabolic growth factors including the PDGF, FGF, EGF, NGF, HGF and Insulin receptor subfamilies. In addition there are a large family of RPTKs for which the corresponding ligand is unknown. Ligands for RPTKs include mainly secreted small proteins, but also membrane-bound and extracellular matrix proteins.

Activation of RPTK by ligands involves ligand-mediated receptor dimerization, resulting in transphosphorylation of the receptor subunits and activation of the cytoplasmic tyrosine kinases. The cytoplasmic tyrosine kinases include receptor associated tyrosine kinases of the src-family (e.g., src, yes, lck, lyn, fyn) and non-receptor linked and cytosolic protein tyrosine kinases, such as the Jak family, members of which mediate signal transduction triggered by the cytokine superfamily of receptors (e.g., the Interleukins, Interferons, GM-CSF, and Leptin).

Because of the wide range of known factors capable of stimulating tyrosine kinase activity, the identification of novel human secreted proteins capable of activating tyrosine kinase signal transduction pathways are of interest. Therefore, the following protocol is designed to identify those novel human secreted proteins capable of activating the tyrosine kinase signal transduction pathways.

Seed target cells (e.g., primary keratinocytes) at a density of approximately 25,000 cells per well in a 96 well Loprodyn Silent Screen Plates purchased from Nalge Nunc (Naperville, IL). The plates are sterilized with two 30 minute rinses with 100% ethanol, rinsed with water and dried overnight. Some plates are coated for 2 hr with 100 ml of cell culture grade type I collagen (50 mg/ml), gelatin (2%) or polylysine (50 mg/ml), all of which can be purchased from Sigma Chemicals (St. Louis, MO) or 10% Matrigel purchased from Becton Dickinson (Bedford, MA), or calf serum, rinsed with PBS and stored at 4°C. Cell growth on these plates is assayed by seeding 5,000 cells/well in growth medium and indirect quantitation of cell number through use of alamarBlue as described by the manufacturer Alamar Biosciences, Inc. (Sacramento, CA) after 48 hr. Falcon plate covers #3071 from Becton Dickinson (Bedford, MA) are used to cover the Loprodyn Silent Screen Plates. Falcon Microtest III cell culture plates can also be used in some proliferation experiments.

To prepare extracts, A431 cells are seeded onto the nylon membranes of Loprodyn plates (20,000/200ml/well) and cultured overnight in complete medium. Cells are quiesced by incubation in serum-free basal medium for 24 hr. After 5-20 minutes treatment with EGF (60ng/ml) or 50 ul of the supernatant produced in Example 11, the medium was removed and 100 ml of extraction buffer ((20 mM HEPES pH 7.5, 0.15 M NaCl, 1% Triton X-100, 0.1% SDS, 2 mM Na₃VO₄, 2 mM Na₄P₂O₇ and a cocktail of protease inhibitors (# 1836170) obtained from Boehringer Mannheim (Indianapolis, IN) is added to each well and the plate is shaken on a rotating shaker for 5 minutes at 4°C. The plate is then placed in a vacuum transfer manifold and the extract filtered through the 0.45 mm membrane bottoms of each well using house vacuum. Extracts are collected in a 96-well catch/assay plate in the bottom of the vacuum manifold and immediately placed on ice. To obtain extracts clarified by centrifugation, the content of each well, after detergent solubilization for 5 minutes, is removed and centrifuged for 15 minutes at 4°C at 16,000 x g.

Test the filtered extracts for levels of tyrosine kinase activity. Although many methods of detecting tyrosine kinase activity are known, one method is described here.

Generally, the tyrosine kinase activity of a supernatant is evaluated by determining its ability to phosphorylate a tyrosine residue on a specific substrate (a biotinylated peptide). Biotinylated peptides that can be used for this purpose include PSK1 (corresponding to amino acids 6-20 of the cell division kinase cdc2-p34) and PSK2 (corresponding to amino acids 1-17 of gastrin). Both peptides are substrates for a range of tyrosine kinases and are available from Boehringer Mannheim.

The tyrosine kinase reaction is set up by adding the following components in order. First, add 10ul of 5uM Biotinylated Peptide, then 10ul ATP/Mg₂⁺ (5mM ATP/50mM MgCl₂), then 10ul of 5x Assay Buffer (40mM imidazole hydrochloride, pH7.3, 40 mM beta-glycerophosphate, 1mM EGTA, 100mM MgCl₂, 5 mM MnCl₂, 0.5 mg/ml BSA), then 5ul of Sodium Vanadate(1mM), and then 5ul of water. Mix the components gently and preincubate the reaction mix at 30°C for 2 min. Initial the reaction by adding 10ul of the control enzyme or the filtered supernatant.

The tyrosine kinase assay reaction is then terminated by adding 10 ul of 120mM EDTA and place the reactions on ice.

Tyrosine kinase activity is determined by transferring 50 ul aliquot of reaction mixture to a microtiter plate (MTP) module and incubating at 37°C for 20 min. This allows the streptavidin coated 96 well plate to associate with the biotinylated peptide. Wash the MTP module with 300ul/well of PBS four times. Next add 75 ul of anti-phosphotyrosine antibody conjugated to horse radish peroxidase(anti-P-Tyr-POD(0.5u/ml)) to each well and incubate at 37°C for one hour. Wash the well as above.

Next add 100ul of peroxidase substrate solution (Boehringer Mannheim) and incubate at room temperature for at least 5 mins (up to 30 min). Measure the absorbance of the sample at 405 nm by using ELISA reader. The level of bound peroxidase activity is quantitated using an ELISA reader and reflects the level of tyrosine kinase activity.

Example 20: High-Throughput Screening Assay Identifying Phosphorylation Activity

As a potential alternative and/or complement to the assay of protein tyrosine kinase activity described in Example 19, an assay which detects activation (phosphorylation) of major intracellular signal transduction intermediates can also be used. For example, as described below one particular assay can detect tyrosine phosphorylation of the Erk-1 and Erk-2 kinases. However, phosphorylation of other molecules, such as Raf, JNK, p38 MAP, Map kinase kinase (MEK), MEK kinase, Src, Muscle specific kinase (MuSK), IRAK, Tec, and Janus, as well as any other phosphoserine, phosphotyrosine, or phosphothreonine molecule, can be detected by substituting these molecules for Erk-1 or Erk-2 in the following assay.

Specifically, assay plates are made by coating the wells of a 96-well ELISA plate with 0.1ml of protein G (1ug/ml) for 2 hr at room temp. (RT). The plates are then rinsed with PBS and blocked with 3% BSA/PBS for 1 hr at RT. The protein G plates are then treated with 2 commercial monoclonal antibodies (100ng/well) against Erk-1

and Erk-2 (1 hr at RT) (Santa Cruz Biotechnology). (To detect other molecules, this step can easily be modified by substituting a monoclonal antibody detecting any of the above described molecules.) After 3-5 rinses with PBS, the plates are stored at 4°C until use.

A431 cells are seeded at 20,000/well in a 96-well Loprodyne filterplate and cultured overnight in growth medium. The cells are then starved for 48 hr in basal medium (DMEM) and then treated with EGF (6ng/well) or 50 ul of the supernatants obtained in Example 11 for 5-20 minutes. The cells are then solubilized and extracts filtered directly into the assay plate.

After incubation with the extract for 1 hr at RT, the wells are again rinsed. As a positive control, a commercial preparation of MAP kinase (10ng/well) is used in place

of A431 extract. Plates are then treated with a commercial polyclonal (rabbit) antibody (1ug/ml) which specifically recognizes the phosphorylated epitope of the Erk-1 and Erk-2 kinases (1 hr at RT). This antibody is biotinylated by standard procedures. The bound polyclonal antibody is then quantitated by successive

incubations with Europium-streptavidin and Europium fluorescence enhancing reagent in the Wallac DELFIA instrument (time-resolved fluorescence). An increased fluorescent signal over background indicates a phosphorylation.

Example 2J: Method of Determining Alterations in a Gene Corresponding to a Polynucleotide

RNA isolated from entire families or individual patients presenting with a phenotype of interest (such as a disease) is isolated. cDNA is then generated from these RNA samples using protocols known in the art. (See, Sambrook.) The cDNA is then used as a template for PCR, employing primers surrounding regions of interest in SEQ ID NO:X. Suggested PCR conditions consist of 35 cycles at 95°C for 30 seconds; 60-120 seconds at 52-58°C; and 60-120 seconds at 70°C, using buffer solutions described in Sidransky, D., et al., Science 252:706 (1991).

PCR products are then sequenced using primers labeled at their 5' end with T4 polynucleotide kinase, employing SequiTherm Polymerase. (Epicentre Technologies). The intron-exon borders of selected exons is also determined and genomic PCR products analyzed to confirm the results. PCR products harboring suspected mutations is then cloned and sequenced to validate the results of the direct sequencing.

PCR products is cloned into T-tailed vectors as described in Holton, T.A. and Graham, M.W., Nucleic Acids Research, 19:1156 (1991) and sequenced with T7 polymerase (United States Biochemical). Affected individuals are identified by mutations not present in unaffected individuals.

Genomic rearrangements are also observed as a method of determining alterations in a gene corresponding to a polynucleotide. Genomic clones isolated according to Example 2 are nick-translated with digoxigenin deoxy-uridine 5'-triphosphate (Boehringer Mannheim), and FISH performed as described in Johnson, Cg. et al., Methods Cell Biol. 35:73-99 (1991). Hybridization with the labeled probe is carried out using a vast excess of human cot-1 DNA for specific hybridization to the corresponding genomic locus.

Chromosomes are counterstained with 4,6-diamino-2-phenylidole and propidium iodide, producing a combination of C- and R-bands. Aligned images for

precise mapping are obtained using a triple-band filter set (Chroma Technology, Brattleboro, VT) in combination with a cooled charge-coupled device camera (Photometrics, Tucson, AZ) and variable excitation wavelength filters. (Johnson, Cv. et al., Genet. Anal. Tech. Appl., 8:75 (1991).) Image collection, analysis and chromosomal fractional length measurements are performed using the ISee Graphical Program System. (Inovision Corporation, Durham, NC.) Chromosome alterations of the genomic region hybridized by the probe are identified as insertions, deletions, and translocations. These alterations are used as a diagnostic marker for an associated disease.

Example 22: Method of Detecting Abnormal Levels of a Polypeptide in a Biological Sample

A polypeptide of the present invention can be detected in a biological sample, and if an increased or decreased level of the polypeptide is detected, this polypeptide is a marker for a particular phenotype. Methods of detection are numerous, and thus, it is understood that one skilled in the art can modify the following assay to fit their particular needs.

For example, antibody-sandwich ELISAs are used to detect polypeptides in a sample, preferably a biological sample. Wells of a microtiter plate are coated with specific antibodies, at a final concentration of 0.2 to 10 ug/ml. The antibodies are either monoclonal or polyclonal and are produced by the method described in Example 10. The wells are blocked so that non-specific binding of the polypeptide to the well is reduced.

The coated wells are then incubated for > 2 hours at RT with a sample containing the polypeptide. Preferably, serial dilutions of the sample should be used to validate results. The plates are then washed three times with deionized or distilled water to remove unbound polypeptide.

Next, 50 ul of specific antibody-alkaline phosphatase conjugate, at a concentration of 25-400 ng, is added and incubated for 2 hours at room temperature. The plates are again washed three times with deionized or distilled water to remove unbound conjugate.

Add 75 μ l of 4-methylumbelliferyl phosphate (MUP) or p-nitrophenyl phosphate (NPP) substrate solution to each well and incubate 1 hour at room temperature. Measure the reaction by a microtiter plate reader. Prepare a standard curve, using serial dilutions of a control sample, and plot polypeptide concentration on the X-axis (log scale) and fluorescence or absorbance of the Y-axis (linear scale). Interpolate the concentration of the polypeptide in the sample using the standard curve.

Example 23: Formulating a Polypeptide

The secreted polypeptide composition will be formulated and dosed in a fashion consistent with good medical practice, taking into account the clinical condition of the individual patient (especially the side effects of treatment with the secreted polypeptide alone), the site of delivery, the method of administration, the scheduling of administration, and other factors known to practitioners. The "effective amount" for purposes herein is thus determined by such considerations.

As a general proposition, the total pharmaceutically effective amount of secreted polypeptide administered parenterally per dose will be in the range of about 1 μ g/kg/day to 10 mg/kg/day of patient body weight, although, as noted above, this will be subject to therapeutic discretion. More preferably, this dose is at least 0.01 mg/kg/day, and most preferably for humans between about 0.01 and 1 mg/kg/day for the hormone. If given continuously, the secreted polypeptide is typically administered at a dose rate of about 1 μ g/kg/hour to about 50 μ g/kg/hour, either by 1-4 injections per day or by continuous subcutaneous infusions, for example, using a mini-pump. An intravenous bag solution may also be employed. The length of treatment needed to observe changes and the interval following treatment for responses to occur appears to vary depending on the desired effect.

Pharmaceutical compositions containing the secreted protein of the invention are administered orally, rectally, parenterally, intracisternally, intravaginally, intraperitoneally, topically (as by powders, ointments, gels, drops or transdermal patch), buccally, or as an oral or nasal spray. "Pharmaceutically acceptable carrier" refers to a non-toxic solid, semisolid or liquid filler, diluent, encapsulating material or formulation auxiliary of any type. The term "parenteral" as used herein refers to

5 modes of administration which include intravenous, intramuscular, intraperitoneal,
10 intrasternal, subcutaneous and intraarticular injection and infusion.

15 The secreted polypeptide is also suitably administered by sustained-release
systems. Suitable examples of sustained-release compositions include semi-
5 permeable polymer matrices in the form of shaped articles, e.g., films, or
microcapsules. Sustained-release matrices include polylactides (U.S. Pat. No.
15 3,773,919, EP 58,481), copolymers of L-glutamic acid and gamma-ethyl-L-glutamate
(Sidman, U. et al., Biopolymers 22:547-556 (1983)), poly (2- hydroxyethyl
methacrylate) (R. Langer, et al., J. Biomed. Mater. Res. 15:167-277 (1981), and R.
20 Langer, Chem. Tech. 12:98-105 (1982)), ethylene vinyl acetate (R. Langer et al.) or
poly-D- (-)-3-hydroxybutyric acid (EP 133,988). Sustained-release compositions
also include liposomally entrapped polypeptides. Liposomes containing the secreted
polypeptide are prepared by methods known per se: DE 3,218,121; Epstein et al.,
25 Proc. Natl. Acad. Sci. USA 82:3688-3692 (1985); Hwang et al., Proc. Natl. Acad. Sci.
USA 77:4030-4034 (1980); EP 52,322; EP 36,676; EP 88,046; EP 143,949; EP
15 142,641; Japanese Pat. Appl. 83-118008; U.S. Pat. Nos. 4,485,045 and 4,544,545; and
EP 102,324. Ordinarily, the liposomes are of the small (about 200-800 Angstroms)
30 unilamellar type in which the lipid content is greater than about 30 mol. percent
cholesterol, the selected proportion being adjusted for the optimal secreted
20 polypeptide therapy.

35 For parenteral administration, in one embodiment, the secreted polypeptide is
formulated generally by mixing it at the desired degree of purity, in a unit dosage
injectable form (solution, suspension, or emulsion), with a pharmaceutically
40 acceptable carrier, i.e., one that is non-toxic to recipients at the dosages and
25 concentrations employed and is compatible with other ingredients of the formulation.
For example, the formulation preferably does not include oxidizing agents and other
compounds that are known to be deleterious to polypeptides.

45 Generally, the formulations are prepared by contacting the polypeptide
uniformly and intimately with liquid carriers or finely divided solid carriers or both.
30 Then, if necessary, the product is shaped into the desired formulation. Preferably the
carrier is a parenteral carrier, more preferably a solution that is isotonic with the blood
50 of the recipient. Examples of such carrier vehicles include water, saline, Ringer's

5 solution, and dextrose solution. Non-aqueous vehicles such as fixed oils and ethyl
oleate are also useful herein, as well as liposomes.

10 The carrier suitably contains minor amounts of additives such as substances
that enhance isotonicity and chemical stability. Such materials are non-toxic to
5 recipients at the dosages and concentrations employed, and include buffers such as
phosphate, citrate, succinate, acetic acid, and other organic acids or their salts;
15 antioxidants such as ascorbic acid; low molecular weight (less than about ten
residues) polypeptides, e.g., polyarginine or tripeptides; proteins, such as serum
albumin, gelatin, or immunoglobulins; hydrophilic polymers such as
20 polyvinylpyrrolidone; amino acids, such as glycine, glutamic acid, aspartic acid, or
arginine; monosaccharides, disaccharides, and other carbohydrates including cellulose
or its derivatives, glucose, manose, or dextrans; chelating agents such as EDTA; sugar
alcohols such as mannitol or sorbitol; counterions such as sodium; and/or nonionic
25 surfactants such as polysorbates, poloxamers, or PEG.

15 The secreted polypeptide is typically formulated in such vehicles at a
concentration of about 0.1 mg/ml to 100 mg/ml, preferably 1-10 mg/ml, at a pH of
about 3 to 8. It will be understood that the use of certain of the foregoing excipients,
30 carriers, or stabilizers will result in the formation of polypeptide salts.

Any polypeptide to be used for therapeutic administration can be sterile.
20 Sterility is readily accomplished by filtration through sterile filtration membranes
(e.g., 0.2 micron membranes). Therapeutic polypeptide compositions generally are
35 placed into a container having a sterile access port, for example, an intravenous
solution bag or vial having a stopper pierceable by a hypodermic injection needle.

40 Polypeptides ordinarily will be stored in unit or multi-dose containers, for
25 example, sealed ampoules or vials, as an aqueous solution or as a lyophilized
formulation for reconstitution. As an example of a lyophilized formulation, 10-ml
vials are filled with 5 ml of sterile-filtered 1% (w/v) aqueous polypeptide solution,
45 and the resulting mixture is lyophilized. The infusion solution is prepared by
reconstituting the lyophilized polypeptide using bacteriostatic Water-for-Injection.

30 The invention also provides a pharmaceutical pack or kit comprising one or
more containers filled with one or more of the ingredients of the pharmaceutical
50 compositions of the invention. Associated with such container(s) can be a notice in

the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration. In addition, the polypeptides of the present invention may be employed in conjunction with other therapeutic compounds.

Example 24: Method of Treating Decreased Levels of the Polypeptide

It will be appreciated that conditions caused by a decrease in the standard or normal expression level of a secreted protein in an individual can be treated by administering the polypeptide of the present invention, preferably in the secreted form. Thus, the invention also provides a method of treatment of an individual in need of an increased level of the polypeptide comprising administering to such an individual a pharmaceutical composition comprising an amount of the polypeptide to increase the activity level of the polypeptide in such an individual.

For example, a patient with decreased levels of a polypeptide receives a daily dose 0.1-100 ug/kg of the polypeptide for six consecutive days. Preferably, the polypeptide is in the secreted form. The exact details of the dosing scheme, based on administration and formulation, are provided in Example 23.

Example 25: Method of Treating Increased Levels of the Polypeptide

Antisense technology is used to inhibit production of a polypeptide of the present invention. This technology is one example of a method of decreasing levels of a polypeptide, preferably a secreted form, due to a variety of etiologies, such as cancer.

For example, a patient diagnosed with abnormally increased levels of a polypeptide is administered intravenously antisense polynucleotides at 0.5, 1.0, 1.5, 2.0 and 3.0 mg/kg day for 21 days. This treatment is repeated after a 7-day rest period if the treatment was well tolerated. The formulation of the antisense polynucleotide is provided in Example 23.

Example 26: Method of Treatment Using Gene Therapy

One method of gene therapy transplants fibroblasts, which are capable of expressing a polypeptide, onto a patient. Generally, fibroblasts are obtained from a subject by skin biopsy. The resulting tissue is placed in tissue-culture medium and separated into small pieces. Small chunks of the tissue are placed on a wet surface of a tissue culture flask, approximately ten pieces are placed in each flask. The flask is turned upside down, closed tight and left at room temperature over night. After 24 hours at room temperature, the flask is inverted and the chunks of tissue remain fixed to the bottom of the flask and fresh media (e.g., Ham's F12 media, with 10% FBS, penicillin and streptomycin) is added. The flasks are then incubated at 37°C for approximately one week.

At this time, fresh media is added and subsequently changed every several days. After an additional two weeks in culture, a monolayer of fibroblasts emerge. The monolayer is trypsinized and scaled into larger flasks.

pMV-7 (Kirschmeier, P.T. et al., DNA, 7:219-25 (1988)), flanked by the long terminal repeats of the Moloney murine sarcoma virus, is digested with EcoRI and HindIII and subsequently treated with calf intestinal phosphatase. The linear vector is fractionated on agarose gel and purified, using glass beads.

The cDNA encoding a polypeptide of the present invention can be amplified using PCR primers which correspond to the 5' and 3' end sequences respectively as set forth in Example 1. Preferably, the 5' primer contains an EcoRI site and the 3' primer includes a HindIII site. Equal quantities of the Moloney murine sarcoma virus linear backbone and the amplified EcoRI and HindIII fragment are added together, in the presence of T4 DNA ligase. The resulting mixture is maintained under conditions appropriate for ligation of the two fragments. The ligation mixture is then used to transform bacteria HB101, which are then plated onto agar containing kanamycin for the purpose of confirming that the vector has the gene of interest properly inserted.

The amphotropic pA317 or GP+am12 packaging cells are grown in tissue culture to confluent density in Dulbecco's Modified Eagles Medium (DMEM) with 10% calf serum (CS), penicillin and streptomycin. The MSV vector containing the gene is then added to the media and the packaging cells transduced with the vector. The packaging cells now produce infectious viral particles containing the gene (the packaging cells are now referred to as producer cells).

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Fresh media is added to the transduced producer cells, and subsequently, the media is harvested from a 10 cm plate of confluent producer cells. The spent media, containing the infectious viral particles, is filtered through a millipore filter to remove detached producer cells and this media is then used to infect fibroblast cells. Media is removed from a sub-confluent plate of fibroblasts and quickly replaced with the media from the producer cells. This media is removed and replaced with fresh media. If the titer of virus is high, then virtually all fibroblasts will be infected and no selection is required. If the titer is very low, then it is necessary to use a retroviral vector that has a selectable marker, such as neo or his. Once the fibroblasts have been efficiently infected, the fibroblasts are analyzed to determine whether protein is produced.

The engineered fibroblasts are then transplanted onto the host, either alone or after having been grown to confluence on cytodex 3 microcarrier beads.

Example 27: Method of Treatment Using Gene Therapy - In Vivo

Another aspect of the present invention is using *in vivo* gene therapy methods to treat disorders, diseases and conditions. The gene therapy method relates to the introduction of naked nucleic acid (DNA, RNA, and antisense DNA or RNA) sequences into an animal to increase or decrease the expression of the polypeptide. The polynucleotide of the present invention may be operatively linked to a promoter or any other genetic elements necessary for the expression of the polypeptide by the target tissue. Such gene therapy and delivery techniques and methods are known in the art, see, for example, WO90/11092, WO98/11779; U.S. Patent NO. 5693622, 5705151, 5580859; Tabata H. et al. (1997) Cardiovasc. Res. 35(3):470-479, Chao J et al. (1997) Pharmacol. Res. 35(6):517-522, Wolff J.A. (1997) Neuromuscul. Disord. 7(5):314-318, Schwartz B. et al. (1996) Gene Ther. 3(5):405-411, Tsurumi Y. et al. (1996) Circulation 94(12):3281-3290 (incorporated herein by reference).

The polynucleotide constructs may be delivered by any method that delivers injectable materials to the cells of an animal, such as, injection into the interstitial space of tissues (heart, muscle, skin, lung, liver, intestine and the like). The

polynucleotide constructs can be delivered in a pharmaceutically acceptable liquid or aqueous carrier.

The term "naked" polynucleotide, DNA or RNA, refers to sequences that are free from any delivery vehicle that acts to assist, promote, or facilitate entry into the cell, including viral sequences, viral particles, liposome formulations, lipofectin or precipitating agents and the like. However, the polynucleotides of the present invention may also be delivered in liposome formulations (such as those taught in Felgner P.L. et al. (1995) *Ann. NY Acad. Sci.* 772:126-139 and Abdallah B. et al. (1995) *Biol. Cell* 85(1):1-7) which can be prepared by methods well known to those skilled in the art.

The polynucleotide vector constructs used in the gene therapy method are preferably constructs that will not integrate into the host genome nor will they contain sequences that allow for replication. Any strong promoter known to those skilled in the art can be used for driving the expression of DNA. Unlike other gene therapies techniques, one major advantage of introducing naked nucleic acid sequences into target cells is the transitory nature of the polynucleotide synthesis in the cells. Studies have shown that non-replicating DNA sequences can be introduced into cells to provide production of the desired polypeptide for periods of up to six months.

The polynucleotide construct can be delivered to the interstitial space of tissues within the an animal, including of muscle, skin, brain, lung, liver, spleen, bone marrow, thymus, heart, lymph, blood, bone, cartilage, pancreas, kidney, gall bladder, stomach, intestine, testis, ovary, uterus, rectum, nervous system, eye, gland, and connective tissue. Interstitial space of the tissues comprises the intercellular fluid, mucopolysaccharide matrix among the reticular fibers of organ tissues, elastic fibers in the walls of vessels or chambers, collagen fibers of fibrous tissues, or that same matrix within connective tissue ensheathing muscle cells or in the lacunae of bone. It is similarly the space occupied by the plasma of the circulation and the lymph fluid of the lymphatic channels. Delivery to the interstitial space of muscle tissue is preferred for the reasons discussed below. They may be conveniently delivered by injection into the tissues comprising these cells. They are preferably delivered to and expressed in persistent, non-dividing cells which are differentiated, although delivery and expression may be achieved in non-differentiated or less completely

5 differentiated cells, such as, for example, stem cells of blood or skin fibroblasts. *In vivo* muscle cells are particularly competent in their ability to take up and express polynucleotides.

10 For the naked polynucleotide injection, an effective dosage amount of DNA or
5 RNA will be in the range of from about 0.05 g/kg body weight to about 50 mg/kg body weight. Preferably the dosage will be from about 0.005 mg/kg to about 20 mg/kg and more preferably from about 0.05 mg/kg to about 5 mg/kg. Of course, as
15 the artisan of ordinary skill will appreciate, this dosage will vary according to the tissue site of injection. The appropriate and effective dosage of nucleic acid sequence
20 can readily be determined by those of ordinary skill in the art and may depend on the condition being treated and the route of administration. The preferred route of administration is by the parenteral route of injection into the interstitial space of tissues. However, other parenteral routes may also be used, such as, inhalation of an aerosol formulation particularly for delivery to lungs or bronchial tissues; throat or
25 mucous membranes of the nose. In addition, naked polynucleotide constructs can be delivered to arteries during angioplasty by the catheter used in the procedure.

30 The dose response effects of injected polynucleotide in muscle *in vivo* is determined as follows. Suitable template DNA for production of mRNA coding for polypeptide of the present invention is prepared in accordance with a standard
20 recombinant DNA methodology. The template DNA, which may be either circular or linear, is either used as naked DNA or complexed with liposomes. The quadriceps muscles of mice are then injected with various amounts of the template DNA.

35 Five to six week old female and male Balb/C mice are anesthetized by intraperitoneal injection with 0.3 ml of 2.5% Avertin. A 1.5 cm incision is made on
40 25 the anterior thigh, and the quadriceps muscle is directly visualized. The template DNA is injected in 0.1 ml of carrier in a 1 cc syringe through a 27 gauge needle over one minute, approximately 0.5 cm from the distal insertion site of the muscle into the knee and about 0.2 cm deep. A suture is placed over the injection site for future
45 localization, and the skin is closed with stainless steel clips.

30 After an appropriate incubation time (e.g., 7 days) muscle extracts are prepared by excising the entire quadriceps. Every fifth 15 um cross-section of the
50 individual quadriceps muscles is histochemically stained for protein expression. A

time course for protein expression may be done in a similar fashion except that quadriceps from different mice are harvested at different times. Persistence of DNA in muscle following injection may be determined by Southern blot analysis after preparing total cellular DNA and HIRT supernatants from injected and control mice.

The results of the above experimentation in mice can be used to extrapolate proper dosages and other treatment parameters in humans and other animals using naked DNA.

Example 28: Transgenic Animals.

The polypeptides of the invention can also be expressed in transgenic animals. Animals of any species, including, but not limited to, mice, rats, rabbits, hamsters, guinea pigs, pigs, micro-pigs, goats, sheep, cows and non-human primates, *e.g.*, baboons, monkeys, and chimpanzees may be used to generate transgenic animals. In a specific embodiment, techniques described herein or otherwise known in the art, are used to express polypeptides of the invention in humans, as part of a gene therapy protocol.

Any technique known in the art may be used to introduce the transgene (*i.e.*, polynucleotides of the invention) into animals to produce the founder lines of transgenic animals. Such techniques include, but are not limited to, pronuclear microinjection (Paterson et al., Appl. Microbiol. Biotechnol. 40:691-698 (1994); Carver et al., Biotechnology (NY) 11:1263-1270 (1993); Wright et al., Biotechnology (NY) 9:830-834 (1991); and Hoppe et al., U.S. Pat. No. 4,873,191 (1989)); retrovirus mediated gene transfer into germ lines (Van der Putten et al., Proc. Natl. Acad. Sci., USA 82:6148-6152 (1985)), blastocysts or embryos; gene targeting in embryonic stem cells (Thompson et al., Cell 56:313-321 (1989)); electroporation of cells or embryos (Lo, 1983, Mol. Cell. Biol. 3:1803-1814 (1983)); introduction of the polynucleotides of the invention using a gene gun (*see, e.g.*, Ulmer et al., Science 259:1745 (1993); introducing nucleic acid constructs into embryonic pluripotent stem cells and transferring the stem cells back into the blastocyst; and sperm-mediated gene transfer (Lavitrano et al., Cell 57:717-723 (1989); etc. For a review of such techniques, *see* Gordon, "Transgenic Animals," Intl. Rev. Cytol. 115:171-229 (1989), which is incorporated by reference herein in its entirety.

Any technique known in the art may be used to produce transgenic clones containing polynucleotides of the invention, for example, nuclear transfer into enucleated oocytes of nuclei from cultured embryonic, fetal, or adult cells induced to quiescence (Campell et al., Nature 380:64-66 (1996); Wilmut et al., Nature 385:810-813 (1997)).

The present invention provides for transgenic animals that carry the transgene in all their cells, as well as animals which carry the transgene in some, but not all their cells, *i.e.*, mosaic animals or chimeric. The transgene may be integrated as a single transgene or as multiple copies such as in concatamers, *e.g.*, head-to-head tandems or head-to-tail tandems. The transgene may also be selectively introduced into, and activated in a particular cell type by following, for example, the teaching of Lasko et al. (Lasko et al., Proc. Natl. Acad. Sci. USA 89:6232-6236 (1992)). The regulatory sequences required for such a cell-type specific activation will depend upon the particular cell type of interest, and will be apparent to those of skill in the art. When it is desired that the polynucleotide transgene be integrated into the chromosomal site of the endogenous gene, gene targeting is preferred. Briefly, when such a technique is to be utilized, vectors containing some nucleotide sequences homologous to the endogenous gene are designed for the purpose of integrating, via homologous recombination with chromosomal sequences, into and disrupting the function of the nucleotide sequence of the endogenous gene. The transgene may also be selectively introduced into a particular cell type, thus inactivating the endogenous gene in only that cell type, by following, for example, the teaching of Gu et al. (Gu et al., Science 265:103-106 (1994)). The regulatory sequences required for such a cell-type specific inactivation will depend upon the particular cell type of interest, and will be apparent to those of skill in the art.

Once transgenic animals have been generated, the expression of the recombinant gene may be assayed utilizing standard techniques. Initial screening may be accomplished by Southern blot analysis or PCR techniques to analyze animal tissues to verify that integration of the transgene has taken place. The level of mRNA expression of the transgene in the tissues of the transgenic animals may also be assessed using techniques which include, but are not limited to, Northern blot analysis of tissue samples obtained from the animal, *in situ* hybridization analysis, and reverse

transcriptase-PCR (rt-PCR). Samples of transgenic gene-expressing tissue may also be evaluated immunocytochemically or immunohistochemically using antibodies specific for the transgene product.

Once the founder animals are produced, they may be bred, inbred, outbred, or crossbred to produce colonies of the particular animal. Examples of such breeding strategies include, but are not limited to: outbreeding of founder animals with more than one integration site in order to establish separate lines; inbreeding of separate lines in order to produce compound transgenics that express the transgene at higher levels because of the effects of additive expression of each transgene; crossing of heterozygous transgenic animals to produce animals homozygous for a given integration site in order to both augment expression and eliminate the need for screening of animals by DNA analysis; crossing of separate homozygous lines to produce compound heterozygous or homozygous lines; and breeding to place the transgene on a distinct background that is appropriate for an experimental model of interest.

Transgenic animals of the invention have uses which include, but are not limited to, animal model systems useful in elaborating the biological function of polypeptides of the present invention, studying conditions and/or disorders associated with aberrant expression, and in screening for compounds effective in ameliorating such conditions and/or disorders.

Example 29: Knock-Out Animals.

Endogenous gene expression can also be reduced by inactivating or "knocking out" the gene and/or its promoter using targeted homologous recombination. (*E.g.*, see Smithies et al., *Nature* 317:230-234 (1985); Thomas & Capecchi, *Cell* 51:503-512 (1987); Thompson et al., *Cell* 5:313-321 (1989); each of which is incorporated by reference herein in its entirety). For example, a mutant, non-functional polynucleotide of the invention (or a completely unrelated DNA sequence) flanked by DNA homologous to the endogenous polynucleotide sequence (either the coding regions or regulatory regions of the gene) can be used, with or without a selectable marker and/or a negative selectable marker, to transfect cells that express polypeptides of the invention *in vivo*. In another embodiment, techniques known in

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the art are used to generate knockouts in cells that contain, but do not express the gene of interest. Insertion of the DNA construct, via targeted homologous recombination, results in inactivation of the targeted gene. Such approaches are particularly suited in research and agricultural fields where modifications to embryonic stem cells can be used to generate animal offspring with an inactive targeted gene (*e.g.*, see Thomas & Capecchi 1987 and Thompson 1989, *supra*). However this approach can be routinely adapted for use in humans provided the recombinant DNA constructs are directly administered or targeted to the required site *in vivo* using appropriate viral vectors that will be apparent to those of skill in the art.

10 In further embodiments of the invention, cells that are genetically engineered to express the polypeptides of the invention, or alternatively, that are genetically engineered not to express the polypeptides of the invention (*e.g.*, knockouts) are administered to a patient *in vivo*. Such cells may be obtained from the patient (*i.e.*, animal, including human) or an MHC compatible donor and can include, but are not limited to fibroblasts, bone marrow cells, blood cells (*e.g.*, lymphocytes), adipocytes, muscle cells, endothelial cells etc. The cells are genetically engineered *in vitro* using recombinant DNA techniques to introduce the coding sequence of polypeptides of the invention into the cells, or alternatively, to disrupt the coding sequence and/or endogenous regulatory sequence associated with the polypeptides of the invention, *e.g.*, by transduction (using viral vectors, and preferably vectors that integrate the transgene into the cell genome) or transfection procedures, including, but not limited to, the use of plasmids, cosmids, YACs, naked DNA, electroporation, liposomes, etc. The coding sequence of the polypeptides of the invention can be placed under the control of a strong constitutive or inducible promoter or promoter/enhancer to achieve expression, and preferably secretion, of the polypeptides of the invention. The engineered cells which express and preferably secrete the polypeptides of the invention can be introduced into the patient systemically, *e.g.*, in the circulation, or intraperitoneally.

45 Alternatively, the cells can be incorporated into a matrix and implanted in the body, *e.g.*, genetically engineered fibroblasts can be implanted as part of a skin graft; genetically engineered endothelial cells can be implanted as part of a lymphatic or vascular graft. (See, for example, Anderson et al. U.S. Patent No. 5,399,349; and

5 Mulligan & Wilson, U.S. Patent No. 5,460,959 each of which is incorporated by reference herein in its entirety).

10 When the cells to be administered are non-autologous or non-MHC compatible cells, they can be administered using well known techniques which
5 prevent the development of a host immune response against the introduced cells. For example, the cells may be introduced in an encapsulated form which, while allowing
15 for an exchange of components with the immediate extracellular environment, does not allow the introduced cells to be recognized by the host immune system.

10 Transgenic and "knock-out" animals of the invention have uses which include, but are not limited to, animal model systems useful in elaborating the biological
20 function of polypeptides of the present invention, studying conditions and/or disorders associated with aberrant expression, and in screening for compounds effective in ameliorating such conditions and/or disorders.

25 It will be clear that the invention may be practiced otherwise than as particularly described in the foregoing description and examples. Numerous
30 modifications and variations of the present invention are possible in light of the above teachings and, therefore, are within the scope of the appended claims.

20 The entire disclosure of each document cited (including patents, patent applications, journal articles, abstracts, laboratory manuals, books, or other
35 disclosures) in the Background of the Invention, Detailed Description, and Examples is hereby incorporated herein by reference. Further, the hard copy of the sequence listing submitted herewith and the corresponding computer readable form are both
40 incorporated herein by reference in their entireties.

Applicant's or agent's file reference number	PZ031PCT	International application No.	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>259</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	
Date of deposit July 27, 1998	Accession Number 203070
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
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Name of depositary institution American Type Culture Collection	
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B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	
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B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
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Address of depositary institution (including postal code and country) 10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	
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(PCT Rule 136ts)

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A. The indications made below relate to the microorganism referred to in the description on page <u>243</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>10801 University Boulevard</u> <u>Manassas, Virginia 20110-2209</u> <u>United States of America</u>	
Date of deposit <u>June 11, 1998</u>	Accession Number <u>209965</u>
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Claims

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What Is Claimed Is:

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1. An isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence at least 95% identical to a sequence selected from the group consisting of:

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(a) a polynucleotide fragment of SEQ ID NO:X or a polynucleotide fragment of the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;

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(b) a polynucleotide encoding a polypeptide fragment of SEQ ID NO:Y or a polypeptide fragment encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;

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(c) a polynucleotide encoding a polypeptide domain of SEQ ID NO:Y or a polypeptide domain encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;

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(d) a polynucleotide encoding a polypeptide epitope of SEQ ID NO:Y or a polypeptide epitope encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X;

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(e) a polynucleotide encoding a polypeptide of SEQ ID NO:Y or the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X, having biological activity;

(f) a polynucleotide which is a variant of SEQ ID NO:X;

(g) a polynucleotide which is an allelic variant of SEQ ID NO:X;

(h) a polynucleotide which encodes a species homologue of the SEQ ID NO:Y;

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(i) a polynucleotide capable of hybridizing under stringent conditions to any one of the polynucleotides specified in (a)-(h), wherein said polynucleotide does not hybridize under stringent conditions to a nucleic acid molecule having a nucleotide sequence of only A residues or of only T residues.

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2. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises a nucleotide sequence encoding a secreted protein.

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3. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises a nucleotide sequence encoding the sequence identified as SEQ ID NO:Y or the polypeptide encoded by the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X.

4. The isolated nucleic acid molecule of claim 1, wherein the polynucleotide fragment comprises the entire nucleotide sequence of SEQ ID NO:X or the cDNA sequence included in ATCC Deposit No:Z, which is hybridizable to SEQ ID NO:X.

5. The isolated nucleic acid molecule of claim 2, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the C-terminus or the N-terminus.

6. The isolated nucleic acid molecule of claim 3, wherein the nucleotide sequence comprises sequential nucleotide deletions from either the C-terminus or the N-terminus.

7. A recombinant vector comprising the isolated nucleic acid molecule of claim 1.

8. A method of making a recombinant host cell comprising the isolated nucleic acid molecule of claim 1.

9. A recombinant host cell produced by the method of claim 8.

10. The recombinant host cell of claim 9 comprising vector sequences.

11. An isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from the group consisting of:

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(a) a polypeptide fragment of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

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(b) a polypeptide fragment of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z, having biological activity;

(c) a polypeptide domain of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

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(d) a polypeptide epitope of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

(e) a secreted form of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

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(f) a full length protein of SEQ ID NO:Y or the encoded sequence included in ATCC Deposit No:Z;

(g) a variant of SEQ ID NO:Y;

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(h) an allelic variant of SEQ ID NO:Y; or

(i) a species homologue of the SEQ ID NO:Y.

12. The isolated polypeptide of claim 11, wherein the secreted form or the full length protein comprises sequential amino acid deletions from either the C-terminus or the N-terminus.

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13. An isolated antibody that binds specifically to the isolated polypeptide of claim 11.

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14. A recombinant host cell that expresses the isolated polypeptide of claim 11.

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15. A method of making an isolated polypeptide comprising:

(a) culturing the recombinant host cell of claim 14 under conditions such that said polypeptide is expressed; and

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(b) recovering said polypeptide.

16. The polypeptide produced by claim 15.

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17. A method for preventing, treating, or ameliorating a medical condition, comprising administering to a mammalian subject a therapeutically effective amount of the polypeptide of claim 11 or the polynucleotide of claim 1.

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18. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:

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(a) determining the presence or absence of a mutation in the polynucleotide of claim 1; and

(b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or absence of said mutation.

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19. A method of diagnosing a pathological condition or a susceptibility to a pathological condition in a subject comprising:

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(a) determining the presence or amount of expression of the polypeptide of claim 11 in a biological sample; and

(b) diagnosing a pathological condition or a susceptibility to a pathological condition based on the presence or amount of expression of the polypeptide.

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20. A method for identifying a binding partner to the polypeptide of claim 11 comprising:

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(a) contacting the polypeptide of claim 11 with a binding partner; and

(b) determining whether the binding partner effects an activity of the polypeptide.

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21. The gene corresponding to the cDNA sequence of SEQ ID NO:Y.

22. A method of identifying an activity in a biological assay, wherein the method comprises:

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(a) expressing SEQ ID NO:X in a cell;

(b) isolating the supernatant;

(c) detecting an activity in a biological assay; and

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(d) identifying the protein in the supernatant having the activity.

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23. The product produced by the method of claim 20.

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<110> Human Genome Sciences, Inc.

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<130> PZ031.PCT

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<213> Homo sapiens

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<212> DNA
<213> Homo sapiens
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<211> 1746

<212> DNA

<213> Homo sapiens

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<211> 2876

<212> DNA

<213> Homo sapiens

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<211> 1052

<212> DNA

<213> Homo sapiens

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 <212> DNA
 <213> Homo sapiens

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<212> DNA

<213> Homo sapiens

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 <212> DNA
 <213> Homo sapiens

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 <212> DNA
 <213> Homo sapiens

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<212> DNA
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<223> n equals a,t,g, or c

<220>
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<213> Homo sapiens

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<210> 43
<211> 1766
<212> DNA
<213> Homo sapiens

<400> 43						
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<210> 44
<211> 2572
<212> DNA
<213> Homo sapiens

<220>
 <221> SITE
 <222> (2527)
 <223> n equals a, t, g, or c

<400> 44
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 catgggcgca cttgtgcgct gcaccaccct gtgcctgggc tactacaaga acattcacga 180
 catcatccct gacagaagtg gcccgagct ggggggagat gcaacaataa gaaagatgct 240
 gagcttctgg tggcctttgg ctctaattct ggccacacag agaatcagtc ggccattgt 300
 caacctcttt gttcccccgg accttggtgg cagttctgca gccacagagg cagtggcgat 360
 tttagacagcc acataccctg tggtcacatg ccatacggct gggtgacgga aatccgtgct 420
 gtgtatccct ctttcgacaa gaataacccc agcaacaaac tgggtgagcac gagcaacaca 480
 gtacagcgag ccacatcaa gaagttcacc ttcgtctgca tggctctgtc actcagctc 540
 tgtttcgtga tgttttgga acccaacgtg tctgagaaaa tcttgataga catcatcgga 600
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<210> 45
 <211> 526
 <212> DNA
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (66)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (106)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (484)

<223> n equals a,t,g, or c

<400> 45

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ctccgccct	ggctggctgg	ccctgctgct	gtgggtctca	gccctgagct	gttctttctc	180
cttgccagct	tcttcccttt	cttctctggt	gccccaaagtc	agaaccagct	acaatttttg	240
aaggactttc	ctcggtcttg	ataaatgcaa	tgcttgcac	gggacatcta	tttgcaagaa	300
gtctcttaaa	gaagaaataa	gatctgacaa	ctggctggct	tcccaccttg	ggactgcctc	360
cggatccct	ttgstttctt	atccttgcaa	attactccar	atgattycca	aaactctggg	420
sccttggtga	ratcttttaa	ctggtcagca	awtwtcaaac	gaaatctcca	aacaggaaat	480
cttntgcttc	ctgcatccac	ccccaaagaa	cttgacatt	gacgtt		526

<210> 46

<211> 1032

<212> DNA

<213> Homo sapiens

<220>

<221> SITE

<222> (974)

<223> n equals a,t,g, or c

<400> 46

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tcagagctgc	gagcatgcgg	gtgctctttg	tggagatttg	ctctgtgcct	gtaggaaggt	180
gggtgtgtgg	tgtgcattgc	agcaacattg	gtggaacaga	tgtgtgtgcc	cccatgctgt	240
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<210> 47

<211> 2680

<212> DNA

<213> Homo sapiens

<400> 47

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aaagtaagcg	attttaactt	tttaattttt	ttaatttttg	ctctgtttct	cacacttgag	180
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<210> 48

<211> 730

<212> DNA

<213> Homo sapiens

<400> 48

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<210> 49

<211> 1275

<212> DNA

<213> Homo sapiens

<400> 49

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catccttctt	tttttggtta	tcttcattca	acgtcacact	gttgtctctc	tctccaagca	180
ccaccccttt	gttccaacta	atggatcaaa	gagttatagc	agcttttaag	gacctgattac	240
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aatgcatagc	aagagggaag	gaaactacag	gagaaaaaaa	gaaaagcccc	caaggatatt	660
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<210> 50

<211> 1762

<212> DNA
<213> Homo sapiens

<220>
<221> SITE
<222> (447)
<223> n equals a.t.g. or c

<400> 50
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<210> 51
<211> 2059
<212> DNA
<213> Homo sapiens

<400> 51
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<210> 52

<211> 3282

<212> DNA

<213> Homo sapiens

<400> 52

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<210> 53

<211> 1860

<212> DNA

<213> Homo sapiens

<400> 53

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 <211> 770
 <212> DNA
 <213> Homo sapiens

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<400> 54
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<210> 55
 <211> 1093
 <212> DNA
 <213> Homo sapiens

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<400> 55
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 <211> 632

<212> DNA
<213> Homo sapiens

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<222> (29)
<223> n equals a,t,g, or c

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<221> SITE
<222> (162)
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<210> 57
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<212> DNA
<213> Homo sapiens

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<222> (1614)
<223> n equals a,t,g, or c

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<210> 58

<211> 619

<212> DNA

<213> Homo sapiens

<220>

<221> SITE

<222> (526)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (619)

<223> n equals a,t,g, or c

<400> 58

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aaaaaaaaa	aaaaaaaaa					619

<210> 59

<211> 1378

<212> DNA

<213> Homo sapiens

<400> 59

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aaattacagg	ctctattaac	agccggaaga	gtgttcctac	tcttcgacgc	ggccgcgaat	1320
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<210> 60

<211> 1126

<212> DNA

<213> Homo sapiens

<220>

<221> SITE

<222> (21)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (35)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (49)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (99)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (1012)

<223> n equals a,t,g, or c

<400> 60

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<210> 61

<211> 2078

<212> DNA

<213> Homo sapiens

<220>

<221> SITE

<222> (337)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (492)

<223> n equals a,t,g, or c

<400> 61

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<210> 62
<211> 762
<212> DNA
<213> Homo sapiens

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<220>
<221> SITE
<222> (10)
<223> n equals a,t,g, or c

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<220>
<221> SITE
<222> (12)
<223> n equals a,t,g, or c

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<220>
<221> SITE
<222> (42)
<223> n equals a,t,g, or c

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<220>
<221> SITE
<222> (219)
<223> n equals a,t,g, or c

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<220>
<221> SITE
<222> (747)
<223> n equals a,t,g, or c

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cttccggctc gtatgttgtg tggaaattgtg agcgatanc aatttcacac aggaaacagc 240

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<210> 63

<211> 1094

<212> DNA

<213> Homo sapiens

<400> 63

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<210> 64

<211> 1361

<212> DNA

<213> Homo sapiens

<400> 64

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<210> 65
<211> 947
<212> DNA
<213> Homo sapiens

<220>
<221> SITE
<222> (67)
<223> n equals a,t,g, or c

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<400> 65
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attagatcca cctggagcac caggtctctc taagtctcac ctgggggaatt cggtccacc 600
tgggtcacca gttccacact agagcactgt gtctgccc agagcacaaa gacctgtctc 660
tcccagact ctctctgact gcagccaggc atagtaccct tgcctgtgtt tgcctcctgg 720
tccacagatt tgggtgctgg gcaggtgccc ggacagtgat gaggtcttgc cgccttaact 780
gtccccccca gtcaactctc ccacaggccc agcaggacgc agtctgagg atcagggatt 840
ctacagctgc attaaatca atcctatcca aaaaaaaaaa aaaaaggcg gccgctctag 900
aggatccaag cttacgtacg cgtgcatgcg acatcaagct ctgaaga 947
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<210> 66
<211> 1376
<212> DNA
<213> Homo sapiens

<220>
<221> SITE
<222> (18)
<223> n equals a,t,g, or c

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<400> 66
ggcggggcgg gcggaagngg cggckgcgcg gccggggcag ccatgtcgcc attgtctgcg 60
gcgcggggcg cctgcgggtt ctacgcggta ggcgcgcggy tgatcctggc gcagctgctg 120
cggcgctgcc gcgggggctt cctggagcca gtttcccccc cagcactga cctgtctgct 180
atagtgcagg gagggacaga tggcattggc tattctacag cgaagcatct ggcgagactt 240
ggcatgcatg ttatcatagc tggaaataat gacagcaaa ccaaacaaat tgtaagcaaa 300
ataaaagaag aaaccttgaa cgacaaagt gaaattttat actgtgactt ggttccatg 360
acttccatcc ggcagtttgt gcagaagttc aagatgaaga agattcctct ccatgtcctg 420
atcaacaatg ctggggtgat gatggtccc cagaggaaaa ccagagatgg attcgaaga 480
catttcggcc tgaactacct agggcacttc ctgctgacca acctctctt ggatacgtg 540
```

```

aaagagtctg ggtccccctg ccacagtgcg aggggtggta cegtctcttc tgccaccat      600
tacgtcgctg agctgaacat ggatgacctt cagagcagtg cctgctactc accccacgca      660
gcctacgccc agagcaagct ggcccttgte ctgttcacct accacctcca gcgggtgctg      720
gcgggtgagg gaagccacgt gaccgccaac gtggtggacc ccgggggtggt caacacggac      780
stctacaagc acgtgttctg ggccaccgt ctggcgaaga agcttctcgg ctggttgctt      840
ttcaagacc ccatgaagg agcgtggact tccatctacg cagcagtcac ccagagctg      900
gaaggagtgt gtggccgtta cctatacaac gagaagaga ccaagtccct ccacgtcacc      960
tacaaccaga aactgcagca gcagctgtgg tctaagagtt gtgagatgac tggggtcctt     1020
gatgtgacct tgtgatatcc tgtctcagga tagctgctgc cccaagaaac acattgcacc     1080
tgccaatagc ttgtgggtct g-gaagactg cgggtgttga gtttctcaca ccacctscc     1140
cacagggctc tgcctcttag ttttgagaca gctgcctcaa cctctgcaga acttcaagaa     1200
gccaaataaa cattttggag gataatcacc ccaagtggtc ttcaaccata aactttgtga     1260
ttccaaagtg cccagtgtgc acagggtgcca taaataatta cattttccaa cataaaaaaa     1320
aaaaaaaaaa aaaaaaaaaa aaaaaaaaaa aaaaaaaaaa aaaaaagggc ggccgc      1376

```

```

<210> 67
<211> 2434
<212> DNA
<213> Homo sapiens

```

```

<220>
<221> SITE
<222> (10)
<223> n equals a,t,g, or c

```

```

<220>
<221> SITE
<222> (12)
<223> n equals a,t,g, or c

```

```

<220>
<221> SITE
<222> (27)
<223> n equals a,t,g, or c

```

```

<220>
<221> SITE
<222> (73)
<223> n equals a,t,g, or c

```

```

<220>
<221> SITE
<222> (75)
<223> n equals a,t,g, or c

```

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<220>
<221> SITE
<222> (103)
<223> n equals a,t,g, or c

```

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<220>
<221> SITE
<222> (130)
<223> n equals a,t,g, or c

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<400> 67
ctgggggtan tncaagaacc ctctgtngga cttagatgic aagctctttc ctttgggcag      60
cgtgtttcct ttctncagat agtgtgctgt gtaactaaa ttngccgggt cgctttccat      120

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ttcctgacan	ttgagatgga	atgccttgac	cattgggtgct	ctgacagaga	agtcattggag	180
tcattgccat	ttcctgggttg	cccttttggg	atgtgatcct	gttagtagag	gttttctagc	240
ttcactaag	atatttcttt	ccctaaccat	catacacttg	gcatgttcca	ttcccacttc	300
ctttcccttc	accttaaaag	agactacccc	tttgcccat	attgccaaac	taattttctc	360
tcgtactctc	tctagtgaat	gatgtgctac	caagcatatg	ccaggctgtg	agaggattat	420
actgagtagt	agaaagaagc	taatttgaag	taaaaattat	ttgtataatt	aagaaagcag	480
attagatgca	catgggtcaac	aggaagttag	ctgtatgtct	gctagttaga	ttcaaaacat	540
cataaagatg	atagcatgtc	aatatattag	cctagccatt	atgttagcct	ttgttaggtg	600
ggcagctttt	ctgctttttc	ccttccctct	tggtgacaac	ggaggaaaac	tccaacagaa	660
atcagtctaa	cagggaattt	gggatcatag	tttatatgca	tctgatttga	aaggagtagt	720
gagggaaggtt	ttcatatcag	atctatcttt	ggatttaaaa	gaacatttat	gaaatcaagc	780
cttctaaccac	tagttataat	tgagaagcaa	cagtaactcc	gtggacagca	atcaagctta	840
aaattgtaaa	taaatatggg	gataattcag	ttgttgcaaa	aaaagggcag	aattcagtag	900
aataaagctc	ttttctctta	cagggtattaa	atgaggacag	agaacctcag	gtgttcttat	960
gctagtgtctt	gctgagtgca	lactaagaaa	gcaattccaa	atagatgtat	acatctagag	1020
agagtgggtat	tagagattca	gtgtatgtat	ttattttacat	gagaggaaaac	tggaatataa	1080
tcccataaat	tattggaata	taatcccata	aattatcacc	ttttatgact	ggaaaatatt	1140
tgccaatgaa	gaaatgggtct	gtaggtattt	gtcttaagat	ttttggcctg	ttaataaaaa	1200
tgtaacttta	acggtttctt	atagttgcct	ttataaagtg	tattgtctaa	aatatttttg	1260
tatcatgtgc	ctttgaaatt	tgacagctga	tttgggtgtt	ggatttctgc	ccagccattt	1320
atcagtattta	tcattttatt	cagtagctgg	caggtgtatt	agacaaacga	gacttaggta	1380
aggaatggaa	cctttcctgt	ggtttgactg	cacatcacac	cagaagactc	cagtatccct	1440
cattccagaa	tgaggaaaaa	gtattctaca	aagaacctaa	tcacctctgt	gaaatctatg	1500
ggatggaaac	agtgtggcct	taggagtcac	atagtctctg	catggtgggg	aggatcatga	1560
tggaatatgt	gaatttctac	ttctagaagt	tgtagaatag	gtcctgcact	ttgcagaat	1620
gtccttcttt	aaacctggct	tattccacag	ctgtagtga	taacatgacc	tggggcttag	1680
ctgctctagc	cctgggttct	tggaagacct	acactgctg	gcccctggcc	atccacctaa	1740
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ttgtgcaaaa	gctttctgtt	taatgcatag	tgttaccgat	ttacatcttg	gttttcagtg	1860
gcactatgtc	taggaggcaa	tatcctttta	aacagtgcct	tggttaagat	agatacttgt	1920
gaatcaaaag	tagcacagaa	atgaactaag	tatatcccat	ttggaattat	attttgtatc	1980
tatttaaaat	ggtttcacct	gttaaagggc	caacagaact	cttggtttta	cttttgtaat	2040
tactgtacag	aaaatttcaa	gagtgtttga	gtgcttgcca	tcagggtgtt	tccttaataa	2100
graggatat	gatcatttac	aggaattata	tatgaaaaaa	gtttttgaaa	tgtatttttg	2160
tgatgtgcta	tggttagggg	aaaccaataa	tttatgattt	taaaacattc	gtatgaaaac	2220
attgtacaat	gtaatatgct	caactttctc	aattttttgc	taatttttct	aagatacatt	2280
aaaaatgttt	tatatttttt	tttaagttaa	atggaccctg	taagaaaatt	aaaaatacca	2340
gaacataaaa	aaaaaaaaaa	aaaaaaaaaa	aaaaaaaaaa	aaaaaaaaaa	aaaaaaaaaa	2400
aaaaaaaaaa	aaaaaaaaaa	aaaaaaaaaa	aaaaa			2434

<210> 68
 <211> 1086
 <212> DNA
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (10)
 <223> n equals a,t,g, or c

<220>
 <221> SITE
 <222> (77)
 <223> n equals a,t,g, or c

<220>
 <221> SITE
 <222> (1056)

<223> n equals a,t,g, or c

<400> 68

ttgaaacacn	cttttgagca	ttaggtccca	gtcccaaccg	ggtggcgggc	gctctagaac	60
tagtggatcc	cccggngtgc	akkaattcgg	cacgagcaac	ctgtagtggag	ttggctgtca	120
ctcagcagct	ctggaaaatt	acctgtctgc	actgaatttc	ctcatcagta	aaatggaaat	180
gattatagta	ctgaccttgt	aggattattc	taaaaatcag	agaagttcat	gcagcttaga	240
acagtgccag	gcacatggca	aatgctatgg	catacttaag	catcttcttc	tgtgtgacct	300
cgctcatacc	atgtgattgt	gccttgcttg	tccctgtttc	acttttcaga	gggagaaaag	360
tggcccaact	taagaatcaa	aattctgatg	ttacttcggg	aaatgcatag	agccagagag	420
acacaatttg	acttagtatg	atccacatca	tcccctcagg	ctgaatagtg	gtggcatgca	480
catctatacc	aaaatgtttt	accttttttg	tagaaggaaa	atatttgtat	cttctatttc	540
atatcttaga	tctttataag	agcacttaag	ttcaacctcc	taagaaactg	ccaattttgt	600
tgatcatgat	agtctgcaca	gattttcgta	ctatttagtg	ktgggagtg	cttagggacc	660
atcaacaaca	ggsccttctt	tttatccatg	agactactga	ggccttgagg	gttattttgt	720
catccatggc	gggtgcacrg	ctagagggtta	tctgggttagt	agcccaacta	agattagaac	780
ccaggaaatt	tgatttaatc	tcgaatgacc	tcttttattt	ctgcattccg	gaaggagaa	840
aggaaagtaa	acgggaaatt	catgttatct	atggaaaagt	tatcagtttg	atgtttatta	900
aatgatttgt	ttcaggagat	tgtttaaca	ttttatcttc	ctagacaata	attttctgta	960
agagtaag	catggtcatt	aaggtagtca	tattaatgta	ttcagtaasc	tgtgaagaaa	1020
aacatatac	aatgtttttc	aataaaatc	agtgantacc	tgaaaaaaa	aaaaaaaaa	1080
aaaaaa						1086

<210> 69

<211> 1262

<212> DNA

<213> Homo sapiens

<220>

<221> SITE

<222> (568)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (639)

<223> n equals a,t,g, or c

<400> 69

cctyctatag	gkmaagctgg	tackcctgca	ggtaccggtc	cggaaattccc	gggtcgaccc	60
acgcgtccgc	ttacaacttc	tctyttgttt	ctgtgtttct	tccacaccat	tatggcgtat	120
ttactgatct	acaaccgggt	ttaagttagt	tgccctgcta	aaattttctc	atgaattctc	180
attacatttt	caaacaaaat	ccaaagtgtt	taccatgacc	ttgtggatat	ctttctgacc	240
tcatttacta	ctgttatctt	ccttattcat	tccattccac	ccacagtggc	cttcttgcaa	300
gtcagagaat	gcaacaaata	tgctcctgcc	ttggggcatt	tgcaactctg	ttcttttggc	360
ctgggtcact	cacctccaca	ttctccattt	tttatgactt	tcttctcata	tttggatctc	420
tgttcaaatg	tcacctctca	aaaaggcctt	caaaactacc	ctatttaaaa	tagcacctgc	480
cactgccatc	tccataccca	ttttcatttt	tccatagcac	ttatccctac	ttggcattat	540
gttatataat	tgtttatctg	kttactgnct	gtctcccca	ctggactgta	agctccatga	600
ctgtaagtac	atagatgaat	tacaaaatga	atcaatggng	aawtatcctt	gmcatatatt	660
attttaatgat	tttgmctcca	tttfaagtaa	aaaaaaaaata	ccattttttt	cactcttcaa	720
agtgatatag	tttaactctc	taaactacat	ttttctcatt	tcccgattta	attaaatcag	780
tgatataaaa	aaaagagtga	tggggataty	tgaagaaga	ctaaaaataga	tgccaggaaa	840
tactaaaact	gctgaagtgt	agtgggtatat	tttttcttta	cacacagtat	taittgagtt	900
actaatgtg	tcactgaatt	acaqaataag	caaaatacta	ggtaaacaga	atcacgcttg	960
ggggctatat	tttgtgtaaa	atttgtgtta	tgcaaaaaata	atattaaata	tttaattact	1020
acagttttgt	tattctcttc	ttatttttag	aaatgatttg	cagctgagtg	aatcaggaag	1080
tgacagtgat	gactgaagaa	atatytagct	ataaataaaa	atttatacag	catgtataat	1140

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ttatatttga ttaacaataa aaattcctaa gactgagggg aatatgtctt aacttttgat 1200
gataaaagaa attaaatttg attcagaaat ttcaaaaaaa aaaaaaaaaa aaggggcgcc 1260
gc 1262
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<210> 70
<211> 1642
<212> DNA
<213> Homo sapiens

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<400> 70
ggcgcgctgc ccggarctgc ctgggttgcg ctgccggcca cgtcccccgc ccgggcctca 60
ggctccttcc tactgtccga gggccaccag gccgcccggg gccctgctgc ccgggatgcg 120
tctgttacta gaggtagag tctaccttcg tctcacatgt gccacaaagg atggcatggc 180
ccgggagtgcc ccaccacgtl ggctttcacc ccttgcaaaag ccgacttcg ccgagcgaca 240
cagtggtcaag ccacagctc tccaaggagg aagatgggtcc aggtggggag catcycccta 300
gcagcagcct ctgatccctt ggccaagcag gagggaaacca ttagcagcct gaggagctgg 360
ctggctggga gccctgggga cggcccagcc ttgctcccag ctcaccaca agatgtggac 420
agctcttttg ctcatattga ttttytctt gtcccttatct gaaagccatg cggcatccaa 480
cgatccacgc aactttgtcc ctaacaaaat gtggaaggga ttagtcaaga ggaatgcac 540
tgtggaaca gttgataata aaacgtctga ggtgtgaacc atggcagcag cttctcctgt 600
cacattgacc aaagggattc ggcagcccam ctcaactcta tggaagtcac aacagaggac 660
acaagcagga cagatgtgag tgaaccagca acttcaggag gtgcagctga tgggttgacc 720
tccattgtct ccacggctgt gccctccagt acgactgcgc cctccattac gactgcggcc 780
tccagtatga ctgtggcctc cagtgtctcc acgactgcag cctccagtac aactgtggcc 840
tccattgtct ccacgactac agcctcctagt atgactgcgc cctccagcac tcccatgaca 900
cttgcactcc ccgcccacac gtccacttgc acagggcgga cccgtccac taccgccact 960
gggcatccat ctctcagcac agcctctgca caagtgccaa agagcagcgc gttgccaaga 1020
acagcaaccc tggccacatt ggcacacagt gctcagactg tagcgaccac agcaaacaca 1080
agcagcccca tgagcactcg tccaagtctt tccaagcaca tgcccagtga caccgcggca 1140
agccctgtac cccctatgck tccccagca caaggtccca ttagccaggt gtcagtggac 1200
cagcctgtgg ttaacacaac awataaatcc acamccatgc cctcaaacac aaccmwag 1260
cccctcacc aggcgtgggt agacaaaact ctccttctgg tgggtgtgtt actcgggggt 1320
acccttttca tcacagtctt ggttttgtt gccctgcagg cctatgagag ctacaagaag 1380
aaggactaca cccaggtgga ctacttaac aacgggatgt atgcggactc agaaatgtga 1440
ggggggcggg ggcctggcgg gaggcctggc cccttctctg tcctttcctt ttgcctttga 1500
gaccaaacca agtgcttcca aattcttttg gtgcaattga ggagatagc cagatgctta 1560
aacacattta attgctgtca gattaattcc atgatcacta aagagctgct gcttttttca 1620
taaaaaaaaa aaaaaaaaaa gg 1642
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<210> 71
<211> 921
<212> DNA
<213> Homo sapiens

<220>
<221> SITE
<222> (4)
<223> n equals a,t,g, or c

<220>
<221> SITE
<222> (9)
<223> n equals a,t,g, or c

<220>
<221> SITE
<222> (11)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (15)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (20)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (901)

<223> n equals a,t,g, or c

<400> 71

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gcggccgctc	tagaactagt	ggatcccccg	ggctgcagga	attcggcacg	aggctctgagc	120
agataagatt	aagggtctgg	tctgtgctca	attaactcct	gtgggcacgg	gggctgggaa	180
gagcaaagtc	agcgggtgcct	acagtcagca	ccatgctggg	cctgcccgtg	aaggggaggtc	240
tgctctgggc	gctgctgctg	cttctcttag	gctcccagat	cctgctgctg	tatgcctggc	300
atttccacga	gcaaagggac	tgtgatgaac	acaatgtcat	ggctcgttac	ctccctgcca	360
cagtggagtt	tgctgtccac	acattcaacc	aacagagcaa	ggactactat	gcctacagac	420
tggggcacat	cttgaattcc	tgggaaggagc	agggtggagtc	caagactgta	ttctcaatgg	480
agctactgct	ggggagaact	agggtgtggga	aatttgaaga	cgacattgac	aactgccatt	540
tccaagaag	cacagagctg	aacaatactt	tcacctgctt	cttcaccatc	agcaccaggc	600
cctggatgac	tcagttcagc	ctcctgaaca	agaactgctt	ggagggattc	cactgagtga	660
aacccactca	caggcttgcc	catgtgctgc	tcccacattc	cgtggacatc	agcactactc	720
tyctgaggac	tcttcagtg	ctgagcagct	ttggacttgt	ttgttatect	attttgcatg	780
tgtttgagat	ctcagatcag	tgttttagaa	aatccacaca	tcttgagcct	aatcatgtag	840
tgtagatcat	taaacatcag	catttttaaga	aaaaaaaaaa	aaaaaaaaarct	cgaggggggg	900
nccggtaccc	agggcggaag	a				921

<210> 72

<211> 906

<212> DNA

<213> Homo sapiens

<220>

<221> SITE

<222> (34)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (833)

<223> n equals a,t,g, or c

<400> 72

ggaaattctc	cctcactaat	tggaaacaaa	gctngagctc	caccgcggtg	gcggccgctc	60
tagaactagt	ggatcccccg	ggctgcagga	attcggcacg	agggaagaga	gaggggaggg	120
tgagcagagg	acagggccgg	agttttccgg	gaacggagga	agagcagtgg	aggctgccag	180
gatgaggctg	ctgtgtggcc	tgtggctgtg	gctctccttg	ctgaaagtcc	tgacggccca	240
gacccaacc	ccccgccac	tcccccccc	gatgcagagc	ttccaaggaa	accagttcca	300
gggggaatgg	ttcgtcctgg	gcctggcggg	caacagcttc	aggccggagc	acagggcgct	360
gctgaacgct	ttcaccgcaa	ctttcgagct	aagtgatgat	ggccgctttg	aggctgtgaa	420

tgcgatgact	cgaggccagc	actgtgacac	atgggtcttat	gtgctgatac	cggcagccca	480
gcctgggag	ttcactgtgg	accacgggtgt	gggcaggagc	tgggtgctgc	ctcccgggac	540
gctggaccag	ttcatctgcc	tgggcagagc	tcarggcctc	tcggatgaca	acattgtctt	600
cccagatgtg	actggargtg	ccctggacct	cacagacctg	ccctgggtgg	cagccccagc	660
ctgaccactc	agacagccgc	ggcccccaag	gcctgactct	tcttgtggga	gggcgaggct	720
ggtcacccca	ggccagcgct	tgttgaagga	tgaagcagct	cctgtccggc	ccagccctgc	780
ctcacagctg	tgcgagctct	gccctcctca	gctctcaaac	ctgaataaat	gcnccaagcc	840
caaaaaaaa	aaaaaaaaa	aaaaaaaaa	ctcgaggggg	ggcccgggtac	ccaattcgga	900
agattg						906

<210> 73
 <211> 680
 <212> DNA
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (7)
 <223> n equals a,t,g, or c

<220>
 <221> SITE
 <222> (9)
 <223> n equals a,t,g, or c

<220>
 <221> SITE
 <222> (15)
 <223> n equals a,t,g, or c

<220>
 <221> SITE
 <222> (16)
 <223> n equals a,t,g, or c

<220>
 <221> SITE
 <222> (22)
 <223> n equals a,t,g, or c

<400> 73	
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ccccccgggc	tycaggaatt
gacctgtggg	ccatgatgct
ctcttctctc	tcaccagggg
gagtcttgca	tccggaacca
tgcgagtgcg	actgcgcgga
tttggccagt	atagagcgtg
gagaaatggc	ttagcatcgc
aagaaaatgt	tcttctagt
cctccctacc	cagagctctg
gaagtgggga	gtgattgaaa
aaactcgagg	gggggcccgg
cnagctccac	cgcggtggcg
cgccctctag	aactagtggg
gacctgctgc	tggtccctct
cttctctctt	ggagctcaag
cccagacctg	gccagacaat
gcagggtttc	ttcaaagaat
gaagttggct	cacctgtctc
tgagtgagg	gcctccacca
tttcaatgaa	aaaaaaaaa
aaaaaaaaa	aaaaaaaaa

<210> 74
 <211> 1633
 <212> DNA

<213> Homo sapiens

<400> 74

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ggagaattga	tcagtagct	ggacagaact	ctagatgtgt	gtgtgtgaga	gaaagagagg	120
cagggagaa	gagggaggag	ttaccccccac	gatgacctcc	aacttccctt	tctgcacct	180
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<211> 1022

<212> DNA

<213> Homo sapiens

<400> 75

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 <212> DNA
 <213> Homo sapiens

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 <211> 312
 <212> DNA
 <213> Homo sapiens

<400> 77
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<210> 78
 <211> 1170
 <212> DNA
 <213> Homo sapiens

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<210> 79

<211> 368

<212> DNA

<213> Homo sapiens

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<222> (5)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (13)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (35)

<223> n equals a,t,g, or c

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<221> SITE

<222> (351)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (367)

<223> n equals a,t,g, or c

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<210> 80

<211> 1088

<212> DNA

<213> Homo sapiens

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<221> SITE
<222> (9)
<223> n equals a,t,g, or c

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<222> (11)
<223> n equals a,t,g, or c

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<210> 81
<211> 1862
<212> DNA
<213> Homo sapiens

<400> 81
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 <212> DNA
 <213> Homo sapiens

<400> 82						
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<211> 2034
<212> DNA
<213> Homo sapiens

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 <212> DNA
 <213> Homo sapiens

<400> 84	
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 <212> DNA
 <213> Homo sapiens

<400> 65

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<211> 3174

<212> DNA

<213> Homo sapiens

<400> 86

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<210> 87

<211> 2780

<212> DNA

<213> Homo sapiens

<220>

<221> SITE

<222> (2760)

<223> n equals a,t,g, or c

<400> 87

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 <212> DNA
 <213> Homo sapiens.

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 <212> DNA

<213> Homo sapiens

<400> 89

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<210> 90

<211> 770

<212> DNA

<213> Homo sapiens

<220>

<221> SITE

<222> (690)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (762)

<223> n equals a,t,g, or c

<400> 90

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<212> DNA
<213> Homo sapiens

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<211> 1722

<212> DNA

<213> Homo sapiens

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<211> 635

<212> DNA

<213> Homo sapiens

<400> 94

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<211> 3798

<212> DNA

<213> Homo sapiens

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<210> 97
<211> 2181
<212> DNA
<213> Homo sapiens

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<220>
<221> SITE
<222> (5)
<223> n equals a,t,g, or c

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<400> 97

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aatttttaaa	cctactgtag	tacaacaagc	caggattgcc	cagaatggaa	ttttgggaga	180
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<210> 98

<211> 1957

<212> DNA

<213> Homo sapiens

<400> 98

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cagcaaaagc gatgctgaga ggggaacag tccagagtcc aacagcagaa cttgggggaa      1020
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ctcttcacag tccctggggg ttgaccagga gccggtcaga gatggacctg gccagatgtc      1860
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<210> 99
 <211> 1112
 <212> DNA
 <213> Homo sapiens

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<400> 99
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gaggcacgag agcatcccac ggtggtagcc agtcataaat cacctgtttt ggatacaaag      540
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gcagcggaag agggccagct caagtctgaa gccgcaggca gccagaccca aggcagcaca      720
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<210> 100
 <211> 887
 <212> DNA
 <213> Homo sapiens

 <220>
 <221> SITE
 <222> (303)
 <223> n equals a,t,g, or c

 <400> 100

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tatggactgt gcccctttct atagtgtgt aatgcttcaa caaagtgct aatagtgtg 180
catgcacatg ttaaaatttc aaactatatt aaagagtggt caattaaaag gaagttatcc 240
tctcactcta aagtcctatt tctctcttc agtctatcat tactactagt ttctagtata 300
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aaccctgtct ctactaaaaa tacaataaaa aaaaaaaa actcgta 887

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<210> 101
<211> 1248
<212> DNA
<213> Homo sapiens

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<400> 101
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<210> 102
<211> 1841
<212> DNA
<213> Homo sapiens

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<400> 102
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ctgatcatct tttgcaaatc tgccagcgca tcggctctat gttggataaa gaaattccac 180
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taattgtgaa gtaatgtaag catgtggaag aaaggtgata atgtaactat aaatcatgct 360
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aatgggtccc	tctgcattaa	aaaagttgat	aaagatatag	atgttaagct	aaaaagcttg	480
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<210> 103

<211> 685

<212> DNA

<213> Homo sapiens

<220>

<221> SITE

<222> (678)

<223> n equals a,t,g, or c

<220>

<221> SITE

<222> (679)

<223> n equals a,t,g, or c

<400> 103

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<210> 104

<211> 1168

<212> DNA

<213> Homo sapiens

<400> 104

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cagatacctg	ctgttttctt	gttggttttc	t-gtttttca	ataaataaaa	ctgagtgtta	1140
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<210> 105

<211> 1175

<212> DNA

<213> Homo sapiens

<220>

<221> SITE

<222> (24)

<223> n equals a,t,g, or c

<400> 105

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aagtttttaa	ggatgaacaa	aaattgacca	gacaaaaggag	ttggagaagt	gggarggtac	540
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<210> 106

<211> 1021
 <212> DNA
 <213> Homo sapiens

<400> 106
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 gsgatgtgcat ttaccaatca gaacagatgt tggctttact gcttgattta gttttctgta 180
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 a 1021

<210> 107
 <211> 830
 <212> DNA
 <213> Homo sapiens

<400> 107
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 cctctgactg taaaaagaag tagcagttcc gaaagcaaga gtccctatg aacacgggaag 180
 aagacattgg caacttttga gtacaacaac tatatttaat agagtaattc aagaacatca 240
 gccagtgaat tttatacaag atagtgaag agaaaaggaa gattaattag gggtagttta 300
 ggaatgccatt aaatagccta gaattagggg agtagtcgtt gaatagaaag gaggccacaa 360
 atttgaggga tataagctaa gaattggtaa gccagaaga aggaaaagggt ttgggcagta 420
 aggataaa-ga ggaacaaaat agagaactca gaagcaatat ctgactgtta tcattggaag 480
 aatttttttg cttgcttgag gctggatatt gaagtggatc aggatacttg agtgactatc 540
 tgatgggctt ttggaactag ctctcaagag gtgaaaatta gctttttttt cttt-tcttt 600
 cttttttttt ttttttgagg caaggtctca ctgttgttga ggctgaacct cctgggctca 660
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<210> 108
 <211> 1301
 <212> DNA
 <213> Homo sapiens

<400> 108
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gagatttata	ccacatacca	cacatagcca	cagaaacatc	atcttgaaat	aaagaagagt	1260
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<210> 109

<211> 1932

<212> DNA

<213> Homo sapiens

<400> 109

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aaaaagggaa	atattaggtt	ggtgcacacg	taattgcggt	ttttgcattg	ttgaaatttg	1860
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<210> 110
<211> 1534
<212> DNA
<213> Homo sapiens

<220>
<221> SITE
<222> (1212)
<223> n equals a,t,g, or c

<400> 110
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aaacatttta tatagacaaa aaaaaaaaaa aaaa 1534

<210> 111
<211> 2871
<212> DNA
<213> Homo sapiens

<220>
<221> SITE
<222> (1234)
<223> n equals a,t,g, or c

<220>
<221> SITE
<222> (1259)
<223> n equals a,t,g, or c

<220>
<221> SITE
<222> (1283)
<223> n equals a,t,g, or c

<220>
<221> SITE
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<223> n equals a,t,g, or c

<220>
<221> SITE
<222> (1287)
<223> n equals a,t,g, or c

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<221> SITE
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<223> n equals a,t,g, or c

<220>
<221> SITE
<222> (1912)
<223> n equals a,t,g, or c

<220>
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<223> n equals a,t,g, or c

<220>
<221> SITE
<222> (1935)
<223> n equals a,t,g, or c

<220>
<221> SITE
<222> (1947)
<223> n equals a,t,g, or c

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<210> 113
<211> 2214
<212> DNA
<213> Homo sapiens

<220>
<221> SITE
<222> (289)
<223> n equals a,t,g, or c
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<210> 114
<211> 3300
<212> DNA
<213> Homo sapiens

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 <213> Homo sapiens

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 <223> n equals a,t,g, or c

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 <211> 2189
 <212> DNA
 <213> Homo sapiens

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<212> DNA

<213> Homo sapiens

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<213> Homo sapiens

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<213> Homo sapiens

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<222> (937)
<223> n equals a,t,g, or c

<220>
<221> SITE
<222> (990)
<223> n equals a,t,g, or c

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<213> Homo sapiens

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<221> SITE

<222> (2293)

<223> n equals a,t,g, or c

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tccttttgag	tttgatgctg	attcacttgc	ctttgacatg	gaaaatgacc	ctgctatggg	1440
tacacacaaa	tccaccaaac	aagtagaatt	gactgcacaa	gatgtgaaag	atgcccactg	1500
gtttta:gac	acccttgga	ttacaaaaga	aaatgtgtatt	ctaatcttc	taacagaaaa	1560


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agaagtaaat attgttttgc caacacagtc cattgttcca agaacttttg tgcttaaaac 1620
aggaatgggt ctgttttttg gtgctatagg ccgcatagat ttcttgacag gaaatcagtc 1680
agcttgggtt acagtcgttg cttccaacat cctccctgtg catatcacct ccttggacag 1740
ggcagacgct ctgtatcaga agcatgcagg tcatacgtta ctccagattc caatgggttg 1800
aaaagaacga atggcrggat ttctcctctt tgttgcgaa gacattatgt taaaagaagg 1860
actgggggca cctgaagcag tggccgacat caagttttcc tctgcagggt gggtttcagt 1920
aacacctaat ttttaaggaca gactgcattc ccgaggctat acacctgaag gaacagtttt 1980
gaccgtccgg cccctctctt tgccatatat tgtaaacatc aaaggacagc gcatcaagaa 2040
aagtgtggcc tataaaacca agaagcctcc ttcccttatg tacaacgtga ggaagaagaa 2100
aggaagata aatgtatgag accgaccttg ttccactccag atatttaactg tattgaacac 2160
aacaaaatac attgaatttg tattaaacat ataacgcata aataaagctc ccattcttac 2220
ccttaaaaaa aaaaaaaaag ggcggccgct ctaggaggatc caagcttacg tacgcgtgca 2280
tgcgacgtca tancctcgtt ataggaactg g 2311

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<210> 121
 <211> 1286
 <212> DNA
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (1284)
 <223> n equals a,t,g, or c

```

<400> 121
gggcgcgcgg gtgaaaggcg cattgatgca gcctgcggcg gcctcggagc gcggcggagc 60
agacgtgac cacgttccct tcctcgggtct cctccgcctc cagctccgcg ctgcccgga 120
ggcgggagcc atgcgacccc agggcccccgc cgcctccccg cagcggctcc gcggcctcct 180
gctgctcctg ctgctgcagc tqcccgccgc gtcgagcgcc tctgagatcc ccaaggggaa 240
gcaaaaggcg cactccggca gagggaggtg gtggacctgt ataatggaat gtgcttaca 300
gggcccagcg gagtgccttg tcgagacggg agccctgggg ccaatggcat tccgggtaca 360
cctgggatcc caggctcgga tggattcaaa ggagaaaagg ggaatgtct gagggaagc 420
tttgaggagt cctggacacc caactacaag cagtgttcat ggagttcatt gaattatggc 480
atagatcttg ggaatttgc ggagtgtaca ttacaaaaga tgcgttcaaa tagtgctcta 540
agagcttttg tcagtggctc acctcggcta aaatgcagaa atgcctgctg ccagcgttgg 600
tatttcacat tcaatggagc tgaatgttca ggacctcttc ccattgaagc tataatttat 660
ttggaccaag gaagccctga aatgaattca acaattaata ttcatcgac ttcttctgtg 720
gaaggacttt gtgaaggaaat tgggtctgga ttagtgatg ttgctatctg ggttggcact 780
tggtcagatt acccaaaagg agatgcttct actggatgga attcagtttc tcgcatcatt 840
attgaagaac taccaaaata aatgctttaa ttttcatttg ctacctcttt ttttattatg 900
ccttggaaat gttcacttaa atgacatttt aaataagttt atgtatacat ctgaatgaaa 960
agcaaaagcta aatatgttta cagaccaaag tgtgatttca cactgttttt aaatctagca 1020
ttattcattt tgcttcaatc aaaagtgggt tcaatatatt ttttagtttg ttagaatact 1080
ttcttcatag tcacattctc tcaacctata atttggaaata ttgtgtggt cttttgtttt 1140
ttctcttagt atagcatttt taaaaaata taaaagctac caatcttctg acaatttcta 1200
aatgttaaga atttttttta tatctgttaa ataaaaatta ttccaacaa aaaaaaaaaa 1260
aaaaaaaaaa aaaaaaaaaa aaanaa 1286

```

<210> 122
 <211> 1380
 <212> DNA
 <213> Homo sapiens

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<400> 122
cagaccgcgc gggcaaacgg actggggcca agacccggga gcgcgggcgc aaaggcacca 60
gggcccgcgc agggcgcgc gcacacggcc ttgggggttc tgcgggcctt cgggtgcgcg 120
tctcgcctct agccatgggg tccgcaacgt tggagatcct gggcctgggt ctgtgcctcg 180

```

```

cgggctgggg gggctctgac ctggcgtgcg ggctgcccac gtggcagggtg accgccttcc 240
tggaccacaa categtgacg gcgcagacca cctggaaggg gctgtggatg tegtgtgtgg 300
tgcagagcac gggcacatgc agtgcaaagt gtacgactcg gtgctgggtc tgagcaccga 360
ggtgcaggcg gcgcggggcg tcaccgtgag cgcctgtctg ctggcgttctg ttgcgtcttt 420
cgtgaccttg gcgggcgcgc agtgcaaccac ctgcgtggcc cggggcccg ccaaggcgcg 480
tgtggccctc acgggaggcg tgctctacat gttttgcggg ctgctggcgc tegtgtccact 540
ctgtcgttcc gccaacattg tcgtccgcga gttttacgac ccgtctgtgc ccgtgtcgca 600
gaagtacgag ctgggcgcac gctgtacatc ggctggggcg ccaccgcgct gctcatggta 660
ggcggctgcc tctgtgtctg cgggccttgg gtctgcaccg gccgtccccg cctcagcttc 720
cccgtgaagt actcagcgcc gcggcgcccc acggccaccg gcgactacga caagaagaac 780
tacgtctgag ggcgttgggc acggcggggc ccctcctgcc agccacgcct gcgaggcggt 840
ggataagcct ggggagcccc gcattggacc cggcttccgc cgggtagcgc ggcgcgcagg 900
ctcctcgaa cgtccggctc tcgccccga cgcggctcct ggatccgctc ctgcctgcgc 960
ccgcagctga ccttctctcg ccactagccc ggccctgccc ttaacagacg gaatgaagtt 1020
tccttttctg tgcgcgcgcg tgtttccata ggcagagcgg gtgtcagact gaggatttct 1080
cttccccctc aagacgcgtg gggccttggc tgctgcctta cttcccagag gctcctgtct 1140
acttcggagg ggcggatgca gggccacggg cccccaccg aagatgtgta cacttggtct 1200
ttactccatc ggcaggggcc gagccaggg accagtgaat tggcctggac ctcccggtct 1260
cactccagca tctccccagg caaggcttgt gggcaccgga gcttgagaga gggcgggagt 1320
gggaaggcta agaattctgt taqtaaatgg tttgaactct caaaaaaaaa aaaaaaaaaa 1380

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<210> 123
<211> 3793
<212> DNA
<213> Homo sapiens

<220>
<221> SITE
<222> (1102)
<223> n equals a,t,g, or c

<220>
<221> SITE
<222> (1132)
<223> n equals a,t,g, or c

<220>
<221> SITE
<222> (1199)
<223> n equals a,t,g, or c

<220>
<221> SITE
<222> (1228)
<223> n equals a,t,g, or c

<220>
<221> SITE
<222> (1229)
<223> n equals a,t,g, or c

<220>
<221> SITE
<222> (1231)
<223> n equals a,t,g, or c

<220>
<221> SITE

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<222> (3176)

<223> n equals a, t, g, or c

<400> 123

ggggcttcat	acaggaaatc	tattgtctgt	tcaagttcca	gagaaaagct	tctgttcgtc	60
caagttacta	accaggctaa	accacataga	cgtgaaggaa	ggggctagaa	ggaagggagt	120
gccccactgt	tgatggggta	agaggatcct	gtactgagaa	gttgaccaga	gagggcttca	180
ccatgcgcac	agttccttct	gtaccagtgt	ggaggaaaag	tactgagtga	agggcgaaaa	240
aagagaaaa	agaaatgctc	tgcccttggg	gaactgctaa	cctaggggta	ctgttgatgt	300
tgactatctt	cttagtggtc	gaagcggagg	gtgctgctca	accaaacaac	tcattaatgc	360
tgcaaaactag	caaggagaat	catgctttag	cttcaagcag	tttatgtatg	gatgaaaaac	420
agattacaca	gaactactcg	aaagtactcg	cagaagttaa	cacttcatgg	cctgtaaaag	480
tggctacaaa	tgctgtgctt	tggtgcccct	ctatgcgatt	aagaaatttg	atcataataa	540
catgggaaat	aatcctgaga	ggccagcctt	cctgcacaaa	agcctacaag	aaagaaacaa	600
atgagaccaa	ggaacccaac	tgtactgatg	agagaataac	ctgggtctcc	agacctgatc	660
agaattcgga	ccttcagatt	cgtaccgtgg	ccatcactca	tgacgggtat	tacagatgca	720
taatgggtaac	acctgatggg	atatttccatc	gtggatatca	cctccaagtg	ttagtacac	780
ctgaagtgcac	cctgtttcaa	aacaggaata	gaactgcagt	atgcaaggca	gttgccaggga	840
agccagctgc	gcatactctc	tgatcccgag	agggcgattg	tgccactaag	caagaatact	900
ggagcaatgg	cacagtgaat	gttaagagta	catgccactg	ggaggtccac	aatgtgtcta	960
ccgtgaactg	ccagctctcc	catttgactg	gcaacaagag	tctgtacata	gagctacttc	1020
ctgttccagg	tgccaaaaaa	tcatacaaat	tatatattcc	atatatcatc	cttactatta	1080
ttattttgac	catcgtggga	ttcatttggg	tggtgaaagt	caatggctgc	aaaaaatata	1140
aatgaataa	accagaatct	actccagtty	ttgaggagga	tgaaatgcag	ccctatgcnt	1200
tttacacaga	gaagaacaat	cctctctnng	ntactacaaa	caaggtgaag	gcactctgag	1260
cattacaag	tgaagtgcac	acagacctcc	atactttata	agttgttgga	ctctagtacc	1320
aagaaacaac	aacaaacgag	atacattata	attactgtct	gattttctta	cagttctaga	1380
atgaagactt	atattgaaat	taggttttcc	aaggttctta	gaagacattt	taatggattc	1440
tcatttcatac	ccttgtataa	ttggaatttt	tgattcttag	ctgctaccag	ctagtctctt	1500
gaagaactga	tgttattaca	aagaaaatac	atgcccatga	ccaaatatte	aaattgtgca	1560
ggacagttaa	taatgaaac	caaatttcc	caagaaataa	ctgaagaagg	agcaagtgtg	1620
aacagttttt	tgtgtatcct	ttcagaatat	tttaattgtac	atatgacatt	tgtatatgcc	1680
tactgttatat	gtgtcaattt	atgtgtcccc	ttacatatat	catgcacctt	atctttgtca	1740
aggcaccagt	gggaacaata	cactgcatta	ctgttctata	catatgaaaa	cctaataata	1800
taagtcttag	agatcatttt	atatcatgac	aagttagact	acctcattct	ttttaattgt	1860
tatatataat	tccattgtat	agttatatca	ttatttaatt	aaaaacaacc	ctaagtatgg	1920
atattttgat	tcttttaagt	tttgtttatt	tcttttaagt	tttgtttgtg	gtataaaca	1980
taccacatag	aatgttttct	gtgcataat	ctctttgttt	ttgagtatat	ctgtaggata	2040
acttcttga	gtggaattgt	caggtcaaag	ggtttgtgca	ttttactatt	gatataatat	2100
ttaaatttgt	tcaaatatat	atgtcaaat	ccctccaaca	ttgttttaatt	gtgcctttcc	2160
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taatcaacca	gattaaatat	gatgtgagat	tataataaga	attatactat	ttataaaaa	2280
tggaattata	tttttggtca	tgtttgtaag	agagtgaatg	cacgtgtgag	aacattagct	2340
tcttctgaac	tcattatata	tccacagagg	tgttgatact	tgatgcttaa	cagttttgca	2400
gatgtgctac	attggaattg	tgtattttta	tggtgtacat	tctattgtga	tatatatttt	2460
gaataattaa	tgtctattga	ccatataagt	ggcgaaaaat	gcaccataga	ggacatgggg	2520
tattttattta	caaaactatga	gctacataat	aagcaagtgg	ccatgggtag	gcatagccct	2580
ccctccata	tttttgtgga	gcaaaatatt	ggcaatgttt	atgtaaatca	ttgttaatat	2640
catgaaatta	tttttaatta	aaaacataag	tctatttgc	ccatagcaga	aaaaacatga	2700
gaagttttt	catcatgata	gaaattgaaa	caaaatata	tcattcttca	atcataccat	2760
ctgagatttt	aaagacagct	attttgtctt	ataagtatat	ttttctccct	ctagacattt	2820
cagttactat	ggattttgtc	ctcaaggya	cytttagtct	aattttggga	tgtaaagcta	2880
atcttaatga	cacttggcac	atgatatttt	gatcaagcca	ttttgacttg	accaaagagc	2940
agtgtccatt	aggtttctgc	atataaatat	taccaagcaa	tgttcacaat	agacattcatt	3000
acactgtcct	tgaaatttat	taattcttca	tccaaccttg	gttgagctga	ggctcatagt	3060
taggttcaag	actatctggt	tcaattattac	tgaaaaacaa	agtaagacag	tactatgctt	3120
acctcttaac	ttgataatgt	caaaccaggc	atgttaaatg	acatcataga	aaagantcca	3180
agataactta	tagaagttaa	atttatattgt	acagaaaaata	attgtatgaa	aatctctact	3240
atggggcttg	aacatgggtg	aacattagaa	tgatataaaa	aatttatatat	attctccaaa	3300

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tccacgctag acctgtcaaa ttagagaatc tagagattag acctggcgtg tcagcaaggt 3360
catccaggaa gcagaggctg agacggagtt aggtgtgatt acctacatag tcgattacat 3420
tttacaata acattttata tgttcattt actgtgcttt ctecccatcc catttcgtat 3480
cttttccttt gctttgctag atttgcatt tttctctctc tttctgtctc tctctcttcc 3540
aatatctcta ataatttgaa agtaattcat cataactaaa tatctattgg gggtatgctt 3600
cacttacaaa cttctgaaaa cggctttact gagatataat tgatatattt aagtgtagag 3660
tttgttaaat ttgtcacata tttaaaatgt ggactttggg aaatgttgac atagt-ttac 3720
atctgtgaaa ccatcagcat aatcaagata ataaacttgt ccatcacccc ccaaaaaaaaa 3780
aaaaaaaaaaa aaa 3793

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<210> 124
 <211> 370
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (370)
 <223> Xaa equals stop translation

<400> 124
 Met Leu Gly Ala Phe Val Trp Pro Ser Leu Leu Leu Ala Ala Ala
 1 5 10 15
 Cys Ile Cys Leu Leu Thr Phe Ile Asn Cys Ala Tyr Val Lys Trp Gly
 20 25 30
 Thr Leu Val Gln Asp Ile Phe Thr Tyr Ala Lys Val Leu Ala Leu Ile
 35 40 45
 Ala Val Ile Val Ala Gly Ile Val Arg Leu Gly Gln Gly Ala Ser Thr
 50 55 60
 His Phe Glu Asn Ser Phe Glu Gly Ser Ser Phe Ala Val Gly Asp Ile
 65 70 75 80
 Ala Leu Ala Leu Tyr Ser Ala Leu Phe Ser Tyr Ser Gly Trp Asp Thr
 85 90 95
 Leu Asn Tyr Val Thr Glu Glu Ile Lys Asn Pro Glu Arg Asn Leu Pro
 100 105 110
 Leu Ser Ile Gly Ile Ser Met Pro Ile Val Thr Ile Ile Tyr Ile Leu
 115 120 125
 Thr Asn Val Ala Tyr Tyr Thr Val Leu Asp Met Arg Asp Ile Leu Ala
 130 135 140
 Ser Asp Ala Val Ala Val Thr Phe Ala Asp Gln Ile Phe Gly Ile Phe
 145 150 155 160
 Asn Trp Ile Ile Pro Leu Ser Val Ala Leu Ser Cys Phe Gly Gly Leu
 165 170 175
 Asn Ala Ser Ile Val Ala Ala Ser Arg Leu Phe Phe Val Gly Ser Arg
 180 185 190
 Glu Gly His Leu Pro Asp Ala Ile Cys Met Ile His Val Glu Arg Phe

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      195              200              205
Thr Pro Val Pro Ser Leu Leu Phe Asn Gly Ile Met Ala Leu Ile Tyr
  210              215              220
Leu Cys Val Glu Asp Ile Phe Gln Leu Ile Asn Tyr Tyr Ser Phe Ser
  225              230              235              240
Tyr Trp Phe Phe Val Gly Leu Ser Ile Val Gly Gln Leu Tyr Leu Arg
              245              250              255
Trp Lys Glu Pro Asp Arg Pro Arg Pro Leu Lys Leu Ser Val Phe Phe
              260              265              270
Pro Ile Val Phe Cys Leu Cys Thr Ile Phe Leu Val Ala Val Pro Leu
  275              280              285
Tyr Ser Asp Thr Ile Asn Ser Leu Ile Gly Ile Ala Ile Ala Leu Ser
  290              295              300
Gly Leu Pro Phe Tyr Phe Leu Ile Ile Arg Val Pro Glu His Lys Arg
  305              310              315              320
Pro Leu Tyr Leu Arg Arg Ser Trp Gly Leu Pro Gln Gly Thr Ser Arg
              325              330              335
Ser Cys Val Cys Gln Leu Leu Gln Lys Trp Ile Trp Lys Met Glu Glu
              340              345              350
Arg Cys Pro Ser Asn Gly Ile Pro Ser Leu Thr Lys His His Leu Glu
  355              360              365
Ser Xaa
  370

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<210> 125
<211> 86
<212> PRT
<213> Homo sapiens

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```

<220>
<221> SITE
<222> (86)
<223> Xaa equals stop translation

```

```

<400> 125
Met Gly Phe Trp Cys Gly Cys Pro Phe Cys Leu Leu Val Val Leu Leu
  1              5              10              15
Thr Asp Arg Thr Leu Ser Cys Arg Ser Val Gly Val Pro Cys Asn Val
              20              25              30
Arg Cys Gln Cys Ala Pro Ala Gly Gly Cys Leu Pro Val Arg Leu Leu
              35              40              45
Ala Gly Gln Gly Ser Gly Thr His Leu Arg Arg Gln Ser Ala Arg Ser
  50              55              60

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Gln Ile Ser Ser Cys Met Leu Gly Glu Pro Leu Leu Ser Ser Lys Leu
 65 70 75 80
 Ser Asp Arg Asp Ile Xaa
 85

<210> 126
 <211> 44
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (44)
 <223> Xaa equals stop translation

<400> 126
 Met Tyr Thr Lys Thr His Lys Phe Lys Phe Tyr Asn Phe Leu Ser Leu
 1 5 10 15
 Trp Ile Trp Lys Ile Phe Phe Leu Leu Phe Phe Ile Leu Ile Val Ala
 20 25 30
 Leu Ala Phe Pro Ile Pro Cys Leu Ser Ile Phe Xaa
 35 40

<210> 127
 <211> 319
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (264)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (303)
 <223> Xaa equals any of the naturally occurring L-amino acids

<400> 127
 Met Asn Thr Asp His Leu Arg Leu Thr Val Pro Asn Gly Ile Gly Ala
 1 5 10 15
 Leu Lys Leu Arg Glu Met Glu His Tyr Phe Ser Gln Gly Leu Ser Val
 20 25 30
 Gln Leu Phe Asn Asp Gly Ser Lys Gly Lys Leu Asn His Leu Cys Gly
 35 40 45
 Ala Asp Phe Val Lys Ser His Gln Lys Pro Pro Gln Gly Met Glu Ile
 50 55 60
 Lys Ser Asn Glu Arg Cys Cys Ser Phe Asp Gly Asp Ala Asp Arg Ile
 65 70 75 80

Val Tyr Tyr Tyr His Asp Ala Asp Gly His Phe His Leu Ile Asp Gly
 85 90 95
 Asp Lys Ile Ala Thr Leu Ile Ser Ser Phe Leu Lys Glu Leu Leu Val
 100 105 110
 Glu Ile Gly Glu Ser Leu Asn Ile Gly Val Val Gln Thr Ala Tyr Ala
 115 120 125
 Asn Gly Ser Ser Thr Arg Tyr Leu Glu Glu Val Met Lys Val Pro Val
 130 135 140
 Tyr Cys Thr Lys Thr Gly Val Lys His Leu His His Lys Ala Gln Glu
 145 150 155 160
 Phe Asp Ile Gly Val Tyr Phe Glu Ala Asn Gly His Gly Thr Ala Leu
 165 170 175
 Phe Ser Thr Ala Val Glu Met Lys Ile Lys Gln Ser Ala Glu Gln Leu
 180 185 190
 Glu Asp Lys Lys Arg Lys Ala Ala Lys Met Leu Glu Asn Ile Ile Asp
 195 200 205
 Leu Phe Asn Gln Ala Ala Gly Asp Ala Ile Ser Asp Met Leu Val Ile
 210 215 220
 Glu Ala Ile Leu Ala Leu Lys Gly Leu Thr Val Gln Gln Trp Asp Ala
 225 230 235 240
 Leu Tyr Thr Asp Leu Pro Asn Arg Gln Leu Lys Val Gln Val Ala Asp
 245 250 255
 Arg Arg Val Ile Ser Thr Thr Xaa Ala Glu Arg Gln Ala Val Thr Pro
 260 265 270
 Pro Gly Leu Gln Glu Ala Ile Asn Asp Leu Val Lys Lys Tyr Lys Leu
 275 280 285
 Ser Arg Ala Phe Val Arg Pro Ser Gly Thr Glu Asp Val Val Xaa Ser
 290 295 300
 Ile Cys Arg Ser Arg Leu Thr Arg Lys Cys Arg Ser Pro Cys Thr
 305 310 315

<210> 128

<211> 46

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (46)

<223> Xaa equals stop translation

<430> 128

Met Asp Met Val Cys Phe Cys Ile Tyr Leu Gly Leu Leu Lys Phe Ile
 1 5 10 15

Ser Ala Ile Phe Cys Ser Phe Ser Glu Glu Val Leu Tyr Ile Ser Phe
 20 25 30
 Val Lys Cys Ile Pro Lys Tyr Phe Val Glu Met Leu Leu Xaa
 35 40 45

<210> 129
 <211> 709
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (189)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (275)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (414)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (438)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (641)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (643)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (696)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (697)
 <223> Xaa equals any of the naturally occurring L-amino acids

<400> 129
 Met Ala Gly Leu Asn Cys Gly Val Ser Ile Ala Leu Leu Gly Val Leu
 1 5 10 15
 Leu Leu Gly Ala Ala Arg Leu Pro Arg Gly Ala Glu Ala Phe Glu Ile
 20 25 30

Ala Leu Pro Arg Glu Ser Asn Ile Thr Val Leu Ile Lys Leu Gly Thr
 35 40 45
 Pro Thr Leu Leu Ala Lys Pro Cys Tyr Ile Val Ile Ser Lys Arg His
 50 55 60
 Ile Thr Met Leu Ser Ile Lys Ser Gly Glu Arg Ile Val Phe Thr Phe
 65 70 75 80
 Ser Cys Gln Ser Pro Glu Asn His Phe Val Ile Glu Ile Gln Lys Asn
 85 90 95
 Ile Asp Cys Met Ser Gly Pro Cys Pro Phe Gly Glu Val Gln Leu Gln
 100 105 110
 Pro Ser Thr Ser Leu Leu Pro Thr Leu Asn Arg Thr Phe Ile Trp Asp
 115 120 125
 Val Lys Ala His Lys Ser Ile Gly Leu Glu Leu Gln Phe Ser Ile Pro
 130 135 140
 Arg Leu Arg Gln Ile Gly Pro Gly Glu Ser Cys Pro Asp Gly Val Thr
 145 150 155 160
 His Ser Ile Ser Gly Arg Ile Asp Ala Thr Val Val Arg Ile Gly Thr
 165 170 175
 Phe Cys Ser Asn Gly Thr Val Ser Arg Ile Lys Met Xaa Glu Gly Val
 180 185 190
 Lys Met Ala Leu His Leu Pro Trp Phe His Pro Arg Asn Val Ser Gly
 195 200 205
 Phe Ser Ile Ala Asn Arg Ser Ser Ile Lys Arg Leu Cys Ile Ile Glu
 210 215 220
 Ser Val Phe Glu Gly Glu Gly Ser Ala Thr Leu Met Ser Ala Asn Tyr
 225 230 235 240
 Pro Glu Gly Phe Pro Glu Asp Glu Leu Met Thr Trp Gln Phe Val Val
 245 250 255
 Pro Ala His Leu Arg Ala Ser Val Ser Phe Leu Asn Phe Asn Leu Ser
 260 265 270
 Asn Cys Xaa Arg Lys Glu Glu Arg Val Glu Tyr Tyr Ile Pro Gly Ser
 275 280 285
 Thr Thr Asn Pro Glu Val Phe Lys Leu Glu Asp Lys Gln Pro Gly Asn
 290 295 300
 Met Ala Gly Asn Phe Asn Leu Ser Leu Gln Gly Cys Asp Gln Asp Ala
 305 310 315 320
 Gln Ser Pro Gly Ile Leu Arg Leu Gln Phe Gln Val Leu Val Gln His
 325 330 335
 Pro Gln Asn Glu Ser Asn Lys Ile Tyr Val Val Asp Leu Ser Asn Glu

340	345	350
Arg Ala Met Ser Leu Thr Ile Glu Pro Arg Pro Val Lys Gln Ser Arg 355 360 365		
Lys Phe Val Pro Gly Cys Phe Val Cys Leu Glu Ser Arg Thr Cys Ser 370 375 380		
Ser Asn Leu Thr Leu Thr Ser Gly Ser Lys His Lys Ile Ser Phe Leu 395 390 395 400		
Cys Asp Asp Leu Thr Arg Leu Trp Met Asn Val Glu Lys Xaa Ile Ser 405 410 415		
Cys Thr Asp His Arg Tyr Cys Gln Arg Lys Ser Tyr Ser Leu Gln Val 420 425 430		
Pro Ser Asp Ile Leu Xaa Leu Pro Val Glu Leu His Asp Phe Ser Trp 435 440 445		
Lys Leu Leu Val Pro Lys Asp Arg Leu Ser Leu Val Leu Val Pro Ala 450 455 460		
Gln Lys Leu Gln Gln His Thr His Glu Lys Pro Cys Asn Thr Ser Phe 465 470 475 480		
Ser Tyr Leu Val Ala Ser Ala Ile Pro Ser Gln Asp Leu Tyr Phe Gly 485 490 495		
Ser Phe Cys Pro Gly Gly Ser Ile Lys Gln Ile Gln Val Lys Gln Asn 500 505 510		
Ile Ser Val Thr Leu Arg Thr Phe Ala Pro Ser Phe Arg Gln Glu Ala 515 520 525		
Ser Arg Gln Gly Leu Thr Val Ser Phe Ile Pro Tyr Phe Lys Glu Glu 530 535 540		
Gly Val Phe Thr Val Thr Pro Asp Thr Lys Ser Lys Val Tyr Leu Arg 545 550 555 560		
Thr Pro Asn Trp Asp Arg Gly Leu Pro Ser Leu Thr Ser Val Ser Trp 565 570 575		
Asn Ile Ser Val Pro Arg Asp Gln Val Ala Cys Leu Thr Phe Phe Lys 580 585 590		
Glu Arg Ser Gly Val Val Cys Gln Thr Gly Arg Ala Phe Met Ile Ile 595 600 605		
Gln Glu Gln Arg Thr Arg Ala Glu Glu Ile Phe Ser Leu Asp Glu Asp 610 615 620		
Val Leu Pro Lys Pro Ser Phe His His His Ser Phe Trp Val Asn Ile 625 630 635 640		
Xaa Asn Xaa Ser Pro Thr Ser Gly Lys Gln Leu Asp Leu Leu Phe Ser 645 650 655		

Val Thr Leu Thr Pro Arg Thr Val Asp Leu Thr Val Ile Leu Ile Ala
660 665 670

Ala Val Gly Gly Gly Val Leu Leu Ser Ala Leu Gly Leu Ile Ile
675 680 685

Cys Cys Val Lys Lys Lys Lys Xaa Xaa Thr Arg Gly Pro Ala Val Gly
690 695 700

Ile Tyr Asn Gly Asn
705

<210> 130

<211> 415

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (415)

<223> Xaa equals stop translation

<400> 130

Met Thr Lys Ala Arg Leu Phe Arg Leu Trp Leu Val Leu Gly Ser Val
1 5 10 15

Phe Met Ile Leu Leu Ile Ile Val Tyr Trp Asp Ser Ala Gly Ala Ala
20 25 30

His Phe Tyr Leu His Thr Ser Phe Ser Arg Pro His Thr Gly Pro Pro
35 40 45

Leu Pro Thr Pro Gly Pro Asp Arg Asp Arg Glu Leu Thr Ala Asp Ser
50 55 60

Asp Val Asp Glu Phe Leu Asp Lys Phe Leu Ser Ala Gly Val Lys Gln
65 70 75 80

Ser Asp Leu Pro Arg Lys Glu Thr Glu Gln Pro Pro Ala Pro Gly Ser
85 90 95

Met Glu Glu Asn Val Arg Gly Tyr Asp Trp Ser Pro Arg Asp Ala Arg
100 105 110

Arg Ser Pro Asp Gln Gly Arg Gln Gln Ala Glu Arg Arg Ser Val Leu
115 120 125

Arg Gly Phe Cys Ala Asn Ser Ser Leu Ala Phe Pro Thr Lys Glu Arg
130 135 140

Ala Phe Asp Asp Ile Pro Asn Ser Glu Leu Ser His Leu Ile Val Asp
145 150 155 160

Asp Arg His Gly Ala Ile Tyr Cys Tyr Val Pro Lys Val Ala Cys Thr
165 170 175

Asn Trp Lys Arg Val Met Ile Val Leu Ser Gly Ser Leu Leu His Arg
180 185 190

Gly Ala Pro Tyr Arg Asp Pro Leu Arg Ile Pro Arg Glu His Val His
 195 200 205
 Asn Ala Ser Ala His Leu Thr Phe Asn Lys Phe Trp Arg Arg Tyr Gly
 210 215 220
 Lys Leu Ser Arg His Leu Met Lys Val Lys Leu Lys Lys Tyr Thr Lys
 225 230 235 240
 Phe Leu Phe Val Arg Asp Pro Phe Val Arg Leu Ile Ser Ala Phe Arg
 245 250 255
 Ser Lys Phe Glu Leu Glu Asn Glu Glu Phe Tyr Arg Lys Phe Ala Val
 260 265 270
 Pro Met Leu Arg Leu Tyr Ala Asn His Thr Ser Leu Pro Ala Ser Ala
 275 280 285
 Arg Glu Ala Phe Arg Ala Gly Leu Lys Val Ser Phe Ala Asn Phe Ile
 290 295 300
 Gln Tyr Leu Leu Asp Pro His Thr Glu Lys Leu Ala Pro Phe Asn Glu
 305 310 315 320
 His Trp Arg Gln Val Tyr Arg Leu Cys His Pro Cys Gln Ile Asp Tyr
 325 330 335
 Asp Phe Val Gly Lys Leu Glu Thr Leu Asp Glu Asp Ala Ala Gln Leu
 340 345 350
 Leu Gln Leu Leu Gln Val Asp Arg Gln Leu Arg Phe Pro Pro Ser Tyr
 355 360 365
 Arg Asn Arg Thr Ala Ser Ser Trp Glu Glu Asp Trp Phe Ala Lys Ile
 370 375 380
 Pro Leu Ala Trp Arg Gln Gln Leu Tyr Lys Leu Tyr Glu Ala Asp Phe
 385 390 395 400
 Val Leu Phe Gly Tyr Pro Lys Pro Glu Asn Leu Leu Arg Asp Xaa
 405 410 415

<210> 131
 <211> 242
 <212> PRT
 <213> Homo sapiens

<400> 131
 Met Gln Leu Gly Ser Val Leu Leu Thr Arg Cys Pro Phe Trp Gly Cys
 1 5 10 15
 Phe Ser Gln Leu Met Leu Tyr Ala Glu Arg Ala Glu Ala Arg Arg Lys
 20 25 30
 Pro Asp Ile Pro Val Pro Tyr Leu Tyr Phe Asp Met Gly Ala Ala Val
 35 40 45

Leu Cys Ala Ser Phe Met Ser Phe Gly Val Lys Arg Arg Trp Phe Ala
 50 55 60
 Leu Gly Ala Ala Leu Gln Leu Ala Ile Ser Thr Tyr Ala Ala Tyr Ile
 65 70 75 80
 Gly Gly Tyr Val His Tyr Gly Asp Trp Leu Lys Val Arg Met Tyr Ser
 85 90 95
 Arg Thr Val Ala Ile Ile Gly Gly Phe Leu Val Leu Ala Ser Gly Ala
 100 105 110
 Gly Glu Leu Tyr Arg Arg Lys Pro Arg Ser Arg Ser Leu Gln Ser Thr
 115 120 125
 Gly Gln Val Phe Leu Gly Ile Tyr Leu Ile Cys Val Ala Tyr Ser Leu
 130 135 140
 Gln His Ser Lys Glu Asp Arg Leu Ala Tyr Leu Asn His Leu Pro Gly
 145 150 155 160
 Gly Glu Leu Met Ile Gln Leu Phe Phe Val Leu Tyr Gly Ile Leu Ala
 155 170 175
 Leu Ala Phe Leu Ser Gly Tyr Tyr Val Thr Leu Ala Ala Gln Ile Leu
 180 185 190
 Ala Val Leu Leu Pro Pro Val Met Leu Leu Ile Asp Gly Asn Val Ala
 195 200 205
 Tyr Trp His Asn Thr Arg Arg Val Glu Phe Trp Asn Gln Met Lys Leu
 210 215 220
 Leu Gly Glu Ser Val Gly Ile Phe Gly Thr Ala Val Ile Leu Ala Thr
 225 230 235 240
 Asp Gly

<210> 132
 <211> 313
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (313)
 <223> Xaa equals stop translation

<400> 132
 Met Glu Ser Leu Tyr Asp Leu Trp Glu Phe Tyr Leu Pro Tyr Leu Tyr
 1 5 10 15
 Ser Cys Ile Ser Leu Met Gly Cys Leu Leu Leu Leu Cys Thr Pro
 20 25 30
 Val Gly Leu Ser Arg Met Phe Thr Val Met Gly His Leu Leu Val Lys
 35 40 45

Pro Thr Ile Leu Glu Asp Leu Asp Glu Gln Ile Tyr Ile Ile Thr Leu
 50 55 60
 Glu Glu Glu Ala Leu Gln Arg Arg Leu Asn Gly Leu Ser Ser Ser Val
 65 70 75 80
 Glu Tyr Asn Ile Met Glu Leu Glu Gln Glu Leu Glu Asn Val Lys Thr
 85 90 95
 Leu Lys Thr Lys Leu Glu Arg Arg Lys Lys Ala Ser Ala Trp Glu Arg
 100 105 110
 Asn Leu Val Tyr Pro Ala Val Met Val Leu Leu Leu Ile Glu Thr Ser
 115 120 125
 Ile Ser Val Leu Leu Val Ala Cys Asn Ile Leu Cys Leu Leu Val Asp
 130 135 140
 Glu Thr Ala Met Pro Lys Gly Thr Arg Gly Pro Gly Ile Gly Asn Ala
 145 150 155 160
 Ser Leu Ser Thr Phe Gly Phe Val Gly Ala Ala Leu Glu Ile Ile Leu
 165 170 175
 Ile Phe Tyr Leu Met Val Ser Ser Val Val Gly Phe Tyr Ser Leu Arg
 180 185 190
 Phe Phe Gly Asn Phe Thr Pro Lys Lys Asp Asp Thr Thr Met Thr Lys
 195 200 205
 Ile Ile Gly Asn Cys Val Ser Ile Leu Val Leu Ser Ser Ala Leu Pro
 210 215 220
 Val Met Ser Arg Thr Leu Gly Ile Thr Arg Phe Asp Leu Leu Gly Asp
 225 230 235 240
 Phe Gly Arg Phe Asn Trp Leu Gly Asn Phe Tyr Ile Val Leu Ser Tyr
 245 250 255
 Asn Leu Leu Phe Ala Ile Val Thr Thr Leu Cys Leu Val Arg Lys Phe
 260 265 270
 Thr Ser Ala Val Arg Glu Glu Leu Phe Lys Ala Leu Gly Leu His Lys
 275 280 285
 Leu His Leu Pro Asn Thr Ser Arg Asp Ser Glu Thr Ala Lys Pro Ser
 290 295 300
 Val Asn Gly His Gln Lys Ala Leu Xaa
 305 310

<210> 133
 <211> 183
 <212> PRT
 <213> Homo sapiens
 <220>

<221> SITE
 <222> (183)
 <223> Xaa equals stop translation

 <400> 133
 Met Met Val Cys Ser Ile Met Met Tyr Phe Leu Leu Gly Ile Thr Leu
 1 5 10 15
 Leu Arg Ser Tyr Met Gln Ser Val Trp Thr Glu Glu Ser Gln Cys Thr
 20 25 30
 Leu Leu Asn Ala Ser Ile Thr Glu Thr Phe Asn Cys Ser Phe Ser Cys
 35 40 45
 Gly Pro Asp Cys Trp Lys Leu Ser Gln Tyr Pro Cys Leu Gln Val Tyr
 50 55 60
 Val Asn Leu Thr Ser Ser Gly Glu Lys Leu Leu Tyr His Thr Glu
 65 70 75 80
 Glu Thr Ile Lys Ile Asn Gln Lys Cys Ser Tyr Ile Pro Lys Cys Gly
 85 90 95
 Lys Asn Phe Glu Glu Ser Met Ser Leu Val Asn Val Val Met Glu Asn
 100 105 110
 Phe Arg Lys Tyr Gln His Phe Ser Cys Tyr Ser Asp Pro Glu Gly Asn
 115 120 125
 Gln Lys Ser Val Ile Leu Thr Lys Leu Tyr Ser Ser Asn Val Leu Phe
 130 135 140
 His Ser Leu Phe Trp Pro Thr Cys Met Met Ala Gly Gly Val Ala Ile
 145 150 155 160
 Val Ala Met Val Lys Leu Thr Gln Tyr Leu Ser Leu Leu Cys Glu Arg
 165 170 175
 Ile Gln Arg Ile Asn Arg Xaa
 180

<210> 134
 <211> 147
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (147)
 <223> Xaa equals stop translation

<400> 134
 Met Trp Lys Leu Trp Arg Ala Glu Glu Gly Ala Ala Ala Leu Gly Gly
 1 5 10 15
 Ala Leu Phe Leu Leu Phe Ala Leu Gly Val Arg Gln Leu Leu Lys
 20 25 30

Gln Arg Arg Pro Met Gly Phe Pro Pro Gly Pro Pro Gly Leu Pro Phe
 35 40 45
 Ile Gly Asn Ile Tyr Ser Leu Ala Ala Ser Ser Glu Leu Pro His Val
 50 55 60
 Tyr Met Arg Lys Gln Ser Gln Val Tyr Gly Glu Val Gln Pro Arg Arg
 65 70 75 80
 Ala Pro Gly Arg Glu Gly Arg Gln Ala Gly Pro Gly Trp Pro Gly Pro
 85 90 95
 Ser Trp Leu Asp Leu Trp Pro Pro Leu Gly Arg Leu Val Gly Thr Ser
 100 105 110
 Pro Cys Ala Gly Cys Pro Leu Arg Asp Thr Arg Phe Pro Gly Leu Glu
 115 120 125
 Gly Arg Ser Pro Arg Arg Arg Ala Pro Leu Gln Gly Glu Pro Arg Pro
 130 135 140
 Cys Arg Xaa
 145

<210> 135
 <211> 122
 <212> PRT
 <213> Homo sapiens

<400> 135
 Met Arg Val Arg Ile Gly Leu Thr Leu Leu Cys Ala Val Leu Leu
 1 5 10 15
 Ser Leu Ala Ser Ala Ser Ser Asp Glu Glu Gly Ser Gln Asp Glu Ser
 20 25 30
 Leu Asp Ser Lys Thr Thr Leu Thr Ser Asp Glu Ser Val Lys Asp His
 35 40 45
 Thr Thr Ala Gly Arg Val Val Ala Gly Gln Ile Phe Leu Asp Ser Glu
 50 55 60
 Glu Ser Glu Leu Glu Ser Ser Ile Gln Glu Glu Glu Asp Ser Leu Lys
 65 70 75 80
 Ser Gln Glu Gly Glu Ser Val Thr Glu Asp Ile Ser Phe Leu Glu Ser
 85 90 95
 Pro Asn Pro Glu Asn Lys Asp Tyr Glu Glu Pro Lys Lys Val Arg Lys
 100 105 110
 Pro Gly Ser Leu Asp Ile Phe Leu Ala Phe
 115 120

<210> 136
 <211> 112
 <212> PRT

<213> Homo sapiens

<400> 136

Met Ala Arg Gly Ser Leu Arg Arg Leu Leu Arg Leu Leu Val Leu Gly
 1 5 10 15

Leu Trp Leu Ala Leu Leu Arg Ser Val Ala Gly Glu Gln Ala Pro Gly
 20 25 30

Thr Ala Pro Cys Ser Arg Gly Ser Ser Trp Ser Ala Asp Leu Asp Lys
 35 40 45

Cys Met Asp Cys Ser Thr Ser Cys Pro Leu Pro Ala Ala Leu Ala His
 50 55 60

Pro Trp Gly Arg Ser Glu Pro Asp Leu Arg Ala Gly Ala Ala Phe Trp
 65 70 75 80

Leu Phe Gly Leu Glu Thr Met Pro Gln Arg Glu Lys Phe Thr Thr Pro
 85 90 95

Ile Glu Glu Thr Gly Gly Glu Gly Cys Pro Ala Val Ala Leu Ile Gln
 100 105 110

<210> 137

<211> 140

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (140)

<223> Xaa equals stop translation

<400> 137

Met Leu Leu Gly Pro Val Pro Ile Leu His Ile Lys Ser Gln Leu Trp
 1 5 10 15

Leu Leu Val Leu Ile Leu Val Val Ser Gly Leu Ser Ala Gly Met Ser
 20 25 30

Ile Ile Pro Thr Phe Pro Glu Ile Leu Ser Cys Ala His Glu Asn Gly
 35 40 45

Phe Glu Glu Gly Leu Ser Thr Leu Gly Leu Val Ser Gly Leu Phe Ser
 50 55 60

Ala Met Trp Ser Ile Gly Ala Phe Met Gly Pro Thr Leu Gly Gly Phe
 65 70 75 80

Leu Tyr Glu Lys Ile Gly Phe Glu Trp Ala Ala Ala Ile Gln Gly Leu
 85 90 95

Trp Ala Leu Ile Ser Gly Leu Ala Met Gly Leu Phe Tyr Leu Leu Glu
 100 105 110

Tyr Ser Arg Arg Lys Arg Ser Lys Ser Gln Asn Ile Leu Ser Thr Glu
115 120 125

Glu Glu Arg Thr Thr Leu Leu Pro Asn Glu Thr Xaa
130 135 140

<210> 138
<211> 404
<212> PRT
<213> Homo sapiens

<400> 138
Met Arg Leu Gln Asp Val Tyr Met Leu Asn Val Lys Gly Leu Ala Arg
1 5 10 15
Gly Val Phe Gln Arg Val Thr Gly Ser Ala Ile Thr Asp Leu Tyr Ser
20 25 30
Pro Lys Arg Leu Phe Ser Leu Thr Gly Asp Asp Cys Phe Gln Val Gly
35 40 45
Lys Val Ala Tyr Asp Met Gly Asp Tyr Tyr His Ala Ile Pro Trp Leu
50 55 60
Glu Glu Ala Val Ser Leu Phe Arg Gly Ser Tyr Gly Glu Trp Lys Thr
65 70 75 80
Glu Asp Glu Ala Ser Leu Glu Asp Ala Leu Asp His Leu Ala Phe Ala
85 90 95
Tyr Phe Arg Ala Gly Asn Val Ser Cys Ala Leu Ser Leu Ser Arg Glu
100 105 110
Phe Leu Leu Tyr Ser Pro Asp Asn Lys Arg Met Ala Arg Asn Val Leu
115 120 125
Lys Tyr Glu Arg Leu Leu Ala Glu Ser Pro Asn His Val Val Ala Glu
130 135 140
Ala Val Ile Gln Arg Pro Asn Ile Pro His Leu Gln Thr Arg Asp Thr
145 150 155 160
Tyr Glu Gly Leu Cys Gln Thr Leu Gly Ser Gln Pro Thr Leu Tyr Gln
165 170 175
Ile Pro Ser Leu Tyr Cys Ser Tyr Glu Thr Asn Ser Asn Ala Tyr Leu
180 185 190
Leu Leu Gln Pro Ile Arg Lys Glu Val Ile His Leu Glu Pro Tyr Ile
195 200 205
Ala Leu Tyr His Asp Phe Val Ser Asp Ser Glu Ala Gln Lys Ile Arg
210 215 220
Glu Leu Ala Glu Pro Trp Leu Gln Arg Ser Val Val Ala Ser Gly Glu
225 230 235 240

Lys	Gln	Leu	Gln	Val	Glu	Tyr	Arg	Ile	Ser	Lys	Ser	Ala	Trp	Leu	Lys	
245					250					255						
Asp	Thr	Val	Asp	Leu	Lys	Leu	Val	Thr	Leu	Asn	His	Arg	Ile	Ala	Ala	
260					265					270						
Leu	Thr	Gly	Leu	Asp	Val	Arg	Pro	Pro	Tyr	Ala	Glu	Tyr	Leu	Gln	Val	
275					280					285						
Val	Asn	Tyr	Gly	Ile	Gly	Gly	His	Tyr	Glu	Pro	His	Phe	Asp	His	Ala	
290					295					300						
Thr	Ser	Pro	Ser	Ser	Pro	Leu	Tyr	Arg	Met	Lys	Ser	Gly	Asn	Arg	Val	
305					310					315					320	
Ala	Thr	Phe	Met	Ile	Tyr	Leu	Ser	Ser	Val	Glu	Ala	Gly	Gly	Ala	Thr	
325					330					335						
Ala	Phe	Ile	Tyr	Ala	Asn	Leu	Ser	Val	Pro	Val	Val	Arg	Asn	Ala	Ala	
340					345					350						
Leu	Phe	Trp	Trp	Asn	Leu	His	Arg	Ser	Gly	Glu	Gly	Asp	Ser	Asp	Thr	
355					360					365						
Leu	His	Ala	Gly	Cys	Pro	Val	Leu	Val	Gly	Asp	Lys	Trp	Val	Ala	Asn	
370					375					380						
Lys	Trp	Ile	His	Glu	Tyr	Gly	Gln	Glu	Phe	Arg	Arg	Pro	Cys	Ser	Ser	
385					390					395					400	
Ser Pro Glu Asp																

<210> 139

<211> 96

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (96)

<223> Xaa equals stop translation

<400> 139

Met	Lys	Ala	Pro	His	Thr	Gly	Val	Leu	His	Leu	Gly	Ser	Val	Trp	Val
1				5					10					15	
Phe	Leu	Gly	Pro	Phe	Leu	Leu	Gly	Val	Gly	Tyr	Thr	Leu	Thr	Phe	Asn
			20					25					30		
Pro	Leu	Ser	Gly	Cys	Met	Ser	Thr	Val	Arg	Trp	Leu	Asn	Ser	Asn	Ile
		35					40					45			
Thr	Ala	Asn	Arg	Thr	Leu	Ser	Arg	Ser	Val	Cys	His	Val	Thr	Pro	Leu
	50					55					60				
His	Arg	Ser	Leu	Ser	Pro	His	Asp	Gly	Glu	Tyr	Leu	Arg	Gln	Met	Leu
65					70					75					80

Leu Asn Ser Ser Ser Arg Ala Gly Glu Ala Gly Ser Trp Gly Tyr Xaa
 85 90 95

<210> 140
 <211> 240
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (240)
 <223> Xaa equals stop translation

<400> 140
 Met Gly Ser Cys Ala Arg Leu Leu Leu Leu Trp Gly Cys Thr Val Val
 1 5 10 15
 Ala Ala Gly Leu Ser Gly Val Ala Gly Val Ser Ser Arg Cys Glu Lys
 20 25 30
 Ala Cys Asn Pro Arg Met Gly Asn Leu Ala Leu Gly Arg Lys Leu Trp
 35 40 45
 Ala Asp Thr Thr Cys Gly Gln Asn Ala Thr Glu Leu Tyr Cys Phe Tyr
 50 55 60
 Ser Glu Asn Thr Asp Leu Thr Cys Arg Gln Pro Lys Cys Asp Lys Cys
 65 70 75 80
 Asn Ala Ala Tyr Pro His Leu Ala His Leu Pro Ser Ala Met Ala Asp
 85 90 95
 Ser Ser Phe Arg Phe Pro Arg Thr Trp Trp Gln Ser Ala Glu Asp Val
 100 105 110
 His Arg Glu Lys Ile Gln Leu Asp Leu Glu Ala Glu Phe Tyr Phe Thr
 115 120 125
 His Leu Ile Val Met Phe Lys Ser Pro Arg Pro Ala Ala Met Val Leu
 130 135 140
 Asp Arg Ser Gln Asp Phe Gly Lys Thr Trp Lys Pro Tyr Lys Tyr Phe
 145 150 155 160
 Ala Thr Asn Cys Ser Ala Thr Phe Gly Leu Glu Asp Asp Val Val Lys
 165 170 175
 Lys Gly Ala Ile Cys Thr Ser Lys Tyr Ser Ser Pro Phe Pro Cys Thr
 180 185 190
 Gly Arg Lys Val Ile Phe Lys Ala Leu Ser Pro Pro Tyr Asp Thr Glu
 195 200 205
 Asn Pro Tyr Ser Ala Lys Val Gln Glu Gln Leu Lys Ile Thr Asn Leu

210 215 220
Pro Arg Ala Ala Ala Glu Thr Thr Val Leu Ser Leu Ser Glu Lys Xaa
225 230 235 240

<210> 141
<211> 54
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (54)
<223> Xaa equals stop translation

<400> 141
Met Met Ile Ser Gly Leu Lys Leu Leu Val Leu Phe Leu Lys Phe Ala
1 5 10 15
Pro Glu Asn Tyr Cys Leu Ser Thr Glu Thr Leu Gln Met Pro Asn Arg
20 25 30
His Leu Arg Leu Ser Lys Ala Thr Cys Tyr Leu Met Lys Cys Leu Leu
35 40 45
Pro Ser Tyr Phe Glu Xaa
50

<210> 142
<211> 67
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (67)
<223> Xaa equals stop translation

<400> 142
Met Arg Ser Leu Ile Ser Ser His Pro Cys Gln His Leu Leu Leu Leu
1 5 10 15
Leu Leu Leu Leu Phe Leu Ile Leu Ala Ile Leu Val Asp Val Lys Trp
20 25 30
Tyr Leu Val Leu Phe Ile Cys Ile Ser Leu Met Thr Ser Asp Val Glu
35 40 45
His Leu Phe Met Cys Leu Leu Ala Ile Arg Ile Ser Ser Trp Arg Asn
50 55 60
Val Tyr Xaa
65

<210> 143
 <211> 108
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (48)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (55)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (58)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (67)
 <223> Xaa equals any of the naturally occurring L-amino acids

<400> 143
 Met Phe Tyr Lys Leu Thr Leu Ile Leu Cys Glu Leu Ser Val Ala Gly
 1 5 10 15
 Val Thr Gln Ala Ala Ser Gln Arg Pro Leu Gln Arg Leu Pro Arg His
 20 25 30
 Ile Cys Ser Gln Arg Asn Pro Pro Gly Arg Cys Leu Leu Lys Ala Xaa
 35 40 45
 Leu Gln Thr Thr Trp Gly Xaa Pro Asp Xaa Gln Phe Pro Gly Cys Pro
 50 55 60
 His Pro Xaa Arg Val Thr Leu Asn Ala Arg Gln Met Gly Asn Gly Lys
 65 70 75 80
 Glu Lys Lys Ala Ala Asp Leu Lys Leu Lys Phe Pro Gln Lys Arg Phe
 85 90 95
 Tyr Leu Ser Ala Phe Ser Glu Arg Ile Lys Ala Phe
 100 105

<210> 144
 <211> 84
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (84)
 <223> Xaa equals stop translation

<400> 144

Met Ala Ser Val Gly Thr Thr Leu Val Ser Pro Leu Leu Cys Leu Leu
 1 5 10 15
 Ile Pro Thr Arg Val Ser Asp Pro Trp Leu Gln Asn Thr Pro Leu His
 20 25 30
 Pro Trp Lys Thr Ile Thr Ile Ile Asp Tyr Tyr Leu Ser Leu Gly Phe
 35 40 45
 Leu Gly Trp Thr Gly Leu Ser Trp Val Val His Phe Gly Ala Ser Ala
 50 55 60
 Val Met Gly Arg Gln Trp Leu Gly Ser Leu Gln Arg Leu Pro Cys Ile
 65 70 75 80
 Ser Gly Ser Xaa

<210> 145

<211> 166

<212> PRT

<213> Homo sapiens

<400> 145

Met Gly Ser Arg Phe Leu Leu Val Leu Leu Ser Gly Leu Thr Val Leu
 1 5 10 15
 Leu Ala Leu Pro Gly Ser Glu Ala Lys Asn Ser Gly Ala Ser Cys Pro
 20 25 30
 Pro Cys Pro Lys Tyr Ala Ser Cys His Asn Ser Thr His Cys Thr Cys
 35 40 45
 Glu Asp Gly Phe Arg Ala Arg Ser Gly Arg Thr Tyr Phe His Asp Ser
 50 55 60
 Ser Glu Lys Cys Glu Asp Ile Asn Glu Cys Glu Thr Gly Leu Ala Lys
 65 70 75 80
 Cys Lys Tyr Lys Ala Tyr Cys Arg Asn Lys Val Gly Gly Tyr Ile Cys
 85 90 95
 Ser Cys Leu Val Lys Tyr Thr Leu Phe Asn Phe Leu Ala Gly Ile Ile
 100 105 110
 Asp Tyr Asp His Pro Asp Cys Tyr Glu Asn Asn Ser Gln Gly Thr Thr
 115 120 125
 Gln Ser Asn Val Asp Ile Trp Val Ser Gly Val Lys Pro Gly Phe Gly
 130 135 140
 Lys Gln Leu Val Arg Ile Thr Met Pro Phe Ser Tyr Pro Asn Ile Asn
 145 150 155 160
 Met Ser Ser Cys Asp Phe
 165

<210> 146
 <211> 70
 <212> PRT
 <213> Homo sapiens

<400> 146
 Met Lys Pro Lys His Leu Glu Trp Cys Leu Ala His Ser Trp Cys Val
 1 5 10 15
 Ile Trp Leu Ser Phe Val Ser Pro Pro Thr Ser His Leu Glu Cys Asp
 20 25 30
 Gly Phe Pro Gly Ser Leu Leu Pro Pro Cys Glu Glu Gly Arg Cys Phe
 35 40 45
 Pro Phe Thr Phe His His His Asp Cys His Gly Cys Ser Pro Leu Gln
 50 55 60
 Ser Ser Pro Gly Gln His
 65 70

<210> 147
 <211> 412
 <212> PRT
 <213> Homo sapiens

<400> 147
 Met Cys Cys Trp Pro Leu Leu Leu Leu Trp Gly Leu Leu Pro Gly Thr
 1 5 10 15
 Ala Ala Gly Gly Ser Gly Arg Thr Tyr Pro His Arg Thr Leu Leu Asp
 20 25 30
 Ser Glu Gly Lys Tyr Trp Leu Gly Trp Ser Gln Arg Gly Ser Gln Ile
 35 40 45
 Ala Phe Arg Leu Gln Val Arg Thr Ala Gly Tyr Val Gly Phe Gly Phe
 50 55 60
 Ser Pro Thr Gly Ala Met Ala Ser Ala Asp Ile Val Val Gly Gly Val
 65 70 75 80
 Ala His Gly Arg Pro Tyr Leu Gln Asp Tyr Phe Thr Asn Ala Asn Arg
 85 90 95
 Glu Leu Lys Lys Asp Ala Gln Gln Asp Tyr His Leu Glu Tyr Ala Met
 100 105 110
 Glu Asn Ser Thr His Thr Ile Ile Glu Phe Thr Arg Glu Leu His Thr
 115 120 125
 Cys Asp Ile Asn Asp Lys Ser Ile Thr Asp Ser Thr Val Arg Val Ile
 130 135 140
 Trp Ala Tyr His His Glu Asp Ala Gly Glu Ala Gly Pro Lys Tyr His
 145 150 155 160

Asp Ser Asn Arg Gly Thr Lys Ser Leu Arg Leu Leu Asn Pro Glu Lys
 165 170 175
 Thr Ser Val Leu Ser Thr Ala Leu Pro Tyr Phe Asp Leu Val Asn Gln
 180 185 190
 Asp Val Pro Ile Pro Asn Lys Asp Thr Thr Tyr Trp Cys Gln Met Phe
 195 200 205
 Lys Ile Pro Val Phe Gln Glu Lys His His Val Ile Lys Val Glu Pro
 210 215 220
 Val Ile Gln Arg Gly His Glu Ser Leu Val His His Ile Leu Leu Tyr
 225 230 235 240
 Gln Cys Ser Asn Asn Phe Asn Asp Ser Val Leu Glu Ser Gly His Glu
 245 250 255
 Cys Tyr His Pro Asn Met Pro Asp Ala Phe Leu Thr Cys Glu Thr Val
 260 265 270
 Ile Phe Ala Trp Ala Ile Gly Gly Glu Gly Phe Ser Tyr Pro Pro His
 275 280 285
 Val Gly Leu Ser Leu Gly Thr Pro Leu Asp Pro His Tyr Val Leu Leu
 290 295 300
 Glu Val His Tyr Asp Asn Pro Thr Tyr Glu Glu Gly Leu Ile Asp Asn
 305 310 315 320
 Ser Gly Leu Arg Leu Phe Tyr Thr Met Asp Ile Arg Lys Tyr Asp Ala
 325 330 335
 Gly Val Ile Glu Ala Gly Leu Trp Val Ser Leu Phe His Thr Ile Pro
 340 345 350
 Pro Gly Met Pro Glu Phe Gln Ser Glu Gly His Cys Thr Leu Glu Cys
 355 360 365
 Leu Glu Glu Leu Trp Lys Pro Lys Ser Gln Val Glu Phe Met Cys Leu
 370 375 380
 Leu Phe Phe Ser Met Leu Thr Trp Leu Ala Glu His Gln Ala Ala Ser
 385 390 395 400
 Phe Ser Lys Arg Glu Gly Asn Glu Ile Thr Cys Leu
 405 410

<210> 148

<211> 85

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (85)

<223> Xaa equals stop translation

<400> 148

Met Asn Val Phe Leu Pro Pro Ala Leu Gly Thr Trp Gly Val Ala Arg
 1 5 10 15
 Phe Phe Pro His Leu Val Pro Glu Arg Trp Cys Leu Val Phe Cys Cys
 20 25 30
 Trp Ile Phe Phe Phe Phe Phe Phe Cys Thr Lys Val Ala Thr Arg
 35 40 45
 Ser Val Leu Gly Asp Gln Ala Gly Leu Gly Val Gly Gly Pro His Leu
 50 55 60
 Pro Leu Pro Gly Ser His Ser Val Ser Val Pro Glu Lys Thr Ile Phe
 65 70 75 80
 Ser Leu Lys Gln Xaa
 85

<210> 149

<211> 154

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (154)

<223> Xaa equals stop translation

<400> 149

Met Gly Arg Leu Pro Leu Leu Arg Arg Val Leu Lys Gly Leu Gln Leu
 1 5 10 15
 Leu Leu Ser Leu Leu Ala Phe Ile Cys Glu Glu Val Val Ser Gln Cys
 20 25 30
 Thr Leu Cys Gly Gly Leu Tyr Phe Phe Glu Phe Val Ser Cys Ser Ala
 35 40 45
 Phe Leu Leu Ser Leu Leu Ile Leu Ile Val Tyr Cys Thr Pro Phe Tyr
 50 55 60
 Glu Arg Val Asp Thr Thr Lys Val Lys Ser Ser Asp Phe Tyr Ile Thr
 65 70 75 80
 Leu Gly Thr Gly Cys Val Phe Leu Leu Ala Ser Ile Ile Phe Val Ser
 85 90 95
 Thr His Asp Arg Thr Ser Ala Glu Ile Ala Ala Ile Val Phe Gly Phe
 100 105 110
 Ile Ala Ser Phe Met Phe Leu Leu Asp Phe Ile Thr Met Leu Tyr Glu
 115 120 125
 Lys Arg Gln Glu Ser Gln Leu Arg Lys Pro Glu Asn Thr Thr Arg Ala
 130 135 140
 Glu Ala Leu Thr Glu Pro Leu Asn Ala Xaa

145

150

<210> 150
 <211> 130
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (130)
 <223> Xaa equals stop translation

<400> 150
 Met Arg Gly His Leu Ala Gly Phe Pro Ala Leu Ser Gly Leu Ala Ser
 1 5 10 15
 Val Cys Leu Trp Ala Thr Phe Ser Ala Gln Leu Pro Gly Pro Val Ala
 20 25 30
 Ala Thr Ser Trp Thr Pro Ala Pro Leu Gly Cys Ser Ala Ala Arg Ser
 35 40 45
 Gly Pro Glu Lys Arg Leu Gly Thr Ala Ala Pro Gly Ser Ala Ala Ser
 50 55 60
 Leu Ala Gln Ala Gly Pro Gly Ala Pro Cys Arg Val Leu Pro Val Asp
 65 70 75 80
 Pro Ala Pro Ala Ala Leu Asn Val Arg Glu Pro Gly Trp Leu Gly Gly
 85 90 95
 Leu Phe Asp Gly Ala Leu Leu Gln Val Leu Leu Asn Phe Leu Arg Lys
 100 105 110
 Ser Thr Asp Val Leu Met Asp Thr Arg Glu Ala Glu Ser Leu Glu Val
 115 120 125
 Glu Xaa
 130

<210> 151
 <211> 62
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (62)
 <223> Xaa equals stop translation

<400> 151
 Met Leu Phe Trp Ala Tyr Pro Ile Cys Val Phe Ile Asp Ser Leu Ser
 1 5 10 15
 Cys Gln Pro Cys Leu Trp Ser Thr Gly Ala Thr Ser His Phe Asn Ser
 20 25 30

Pro Thr Thr Ser Pro Leu Phe Thr Leu Phe Met Pro Cys Ala Leu Ala
 35 40 45
 Pro Asn Pro Phe Thr Gln Leu Gly Lys Leu Asp Asp Arg Xaa
 50 55 60

<210> 152
 <211> 225
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (225)
 <223> Xaa equals stop translation

<400> 152
 Met Gly Ile Phe Pro Gly Ile Ile Leu Ile Phe Leu Arg Val Lys Phe
 1 5 10 15
 Ala Thr Ala Ala Val Ile Val Ser Gly His Gln Lys Ser Thr Thr Val
 20 25 30
 Ser His Glu Met Ser Gly Leu Asn Trp Lys Pro Phe Val Tyr Gly Gly
 35 40 45
 Leu Ala Ser Ile Val Ala Glu Phe Gly Thr Phe Pro Val Asp Leu Thr
 50 55 60
 Lys Thr Arg Leu Gln Val Gln Gly Gln Ser Ile Asp Ala Arg Phe Lys
 65 70 75 80
 Glu Ile Lys Tyr Arg Gly Met Phe His Ala Leu Phe Arg Ile Cys Lys
 85 90 95
 Glu Glu Gly Val Leu Ala Leu Tyr Ser Gly Ile Ala Pro Ala Leu Leu
 100 105 110
 Arg Gln Ala Ser Tyr Gly Thr Ile Lys Ile Gly Ile Tyr Gln Ser Leu
 115 120 125
 Lys Arg Leu Phe Val Glu Arg Leu Glu Asp Glu Thr Leu Leu Ile Asn
 130 135 140
 Met Ile Cys Gly Val Val Ser Gly Val Ile Ser Ser Thr Ile Ala Asn
 145 150 155 160
 Pro Thr Asp Val Leu Lys Ile Arg Met Gln Ala Gln Gly Ser Leu Phe
 165 170 175
 Gln Gly Ser Met Ile Gly Ser Phe Ile Asp Ile Tyr Gln Gln Glu Gly
 180 185 190
 Thr Arg Gly Leu Trp Arg Val Ser Thr Leu Phe Leu Leu Ser Tyr
 195 200 205
 Thr Leu Ser Ser Tyr Asn Leu Gln Arg Ile Phe Phe Tyr Ile Lys Thr
 210 215 220

Xaa
225

<210> 153
<211> 69
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (69)
<223> Xaa equals stop translation

<400> 153
Met Leu Met Leu Leu Thr Leu Leu Val Leu Gly Met Val Trp Val Ala
1 5 10 15
Ser Ala Ile Val Asp Lys Asn Lys Ala Asn Arg Glu Ser Leu Tyr Asp
20 25 30
Phe Trp Glu Tyr Tyr Leu Pro Tyr Leu Tyr Ser Cys Ile Ser Phe Leu
35 40 45
Gly Val Leu Leu Leu Leu Ala Ala Gly Arg Pro Gly Gly Ala Ala Val
50 55 60
Leu Leu Ser Leu Xaa
65

<210> 154
<211> 84
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (84)
<223> Xaa equals stop translation

<400> 154
Met Tyr Gly Val Cys Leu Cys Val Ile Val Cys Val Ser Gly Val Ser
1 5 10 15
Leu Cys Leu Tyr Val Trp Gly Val Ser Val Cys Asp Cys Val Ser Val
20 25 30
Phe Met Cys Val Cys Leu Cys Val Ile Phe Cys Val Tyr Gly Lys Pro
35 40 45
Arg Thr Glu His Tyr His Ser Pro His Leu Ala Lys Gln Lys Ala Phe
50 55 60
Arg Glu Met Cys Gly Arg His Asp Val Ser Ala Ala Gly Ile Phe Gln
65 70 75 80
Ser Tyr Val Xaa

<210> 155
<211> 61
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (61)
<223> Xaa equals stop translation

<400> 155
Met His Val Leu Leu Phe Ser Phe Leu Ile Pro Phe Leu Leu Ser
1 5 10 15
Pro Val Gly Val Thr Cys Asn Ser His Met Leu Glu Arg Gln Val Ser
20 25 30
Trp Leu Lys Lys Arg Ser Thr Gln Ala Ser Gln Gln Phe Asn Lys Phe
35 40 45
Leu Arg Gly Ile Ser Asn Val Gly Arg Ile Val Ile Xaa
50 55 60

<210> 156
<211> 84
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (84)
<223> Xaa equals stop translation

<400> 156
Met Cys Leu Leu Val Glu Tyr Ser Leu Met Ile Leu Thr Ile Ile Pro
1 5 10 15
Ser Leu Leu Ser Phe Val Leu Cys Leu Lys Gly Ile Lys His Gly Asn
20 25 30
Tyr Ile Phe Gln Thr Pro Leu Pro Glu Gly Tyr Gly Trp Ile Ser Ala
35 40 45
Met Ser Gly Leu Cys Ile Lys Phe Gly Arg Arg Lys Arg Arg Lys Thr
50 55 60
Trp Leu Leu Gln Val Gly Thr Leu Ala Thr Ile Asp Thr Glu Phe Ala
65 70 75 80
Arg Ser Cys Xaa

<210> 157
<211> 162

<212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (162)
 <223> Xaa equals stop translation

<400> 157
 Met Ala Leu Ser Leu Thr Leu Cys Phe Val Met Phe Trp Thr Pro Asn
 1 5 10 15
 Val Ser Glu Lys Ile Leu Ile Asp Ile Ile Gly Val Asp Phe Ala Phe
 20 25 30
 Ala Glu Leu Cys Val Val Pro Leu Arg Ile Phe Ser Phe Phe Pro Val
 35 40 45
 Pro Val Thr Val Arg Ala His Leu Thr Gly Trp Leu Met Thr Leu Lys
 50 55 60
 Lys Thr Phe Val Leu Ala Pro Ser Ser Val Leu Arg Ile Ile Val Leu
 65 70 75 80
 Ile Ala Ser Leu Val Val Leu Pro Tyr Leu Gly Val His Gly Ala Thr
 85 90 95
 Leu Gly Val Gly Ser Leu Leu Ala Gly Phe Val Gly Glu Ser Thr Met
 100 105 110
 Val Ala Ile Ala Ala Cys Tyr Val Tyr Arg Lys Gln Lys Lys Lys Met
 115 120 125
 Glu Asn Glu Ser Ala Thr Glu Gly Glu Asp Ser Ala Met Thr Asp Met
 130 135 140
 Pro Pro Thr Glu Glu Val Thr Asp Ile Val Glu Met Arg Glu Glu Asn
 145 150 155 160
 Glu Xaa

<210> 158
 <211> 146
 <212> PRT
 <213> Homo sapiens
 <220>
 <221> SITE
 <222> (96)
 <223> Xaa equals any of the naturally occurring L-amino acids
 <220>
 <221> SITE
 <222> (107)
 <223> Xaa equals any of the naturally occurring L-amino acids
 <220>

<221> SITE
 <222> (111)
 <223> Xaa equals any of the naturally occurring L-amino acids

 <220>
 <221> SITE
 <222> (115)
 <223> Xaa equals any of the naturally occurring L-amino acids

 <220>
 <221> SITE
 <222> (122)
 <223> Xaa equals any of the naturally occurring L-amino acids

 <220>
 <221> SITE
 <222> (132)
 <223> Xaa equals any of the naturally occurring L-amino acids

 <400> 158
 Met Glu Pro Gln Leu Gly Pro Glu Ala Ala Leu Arg Pro Gly Trp
 1 5 10 15
 Leu Ala Leu Leu Trp Val Ser Ala Leu Ser Cys Ser Phe Ser Leu
 20 25 30
 Pro Ala Ser Ser Leu Ser Ser Leu Val Pro Gln Val Arg Thr Ser Tyr
 35 40 45
 Asn Phe Gly Arg Thr Phe Leu Gly Leu Asp Lys Cys Asn Ala Cys Ile
 50 55 60
 Gly Thr Ser Ile Cys Lys Lys Phe Phe Lys Glu Glu Ile Arg Ser Asp
 65 70 75 80
 Asn Trp Leu Ala Ser His Leu Gly Thr Ala Ser Arg Phe Pro Leu Xaa
 85 90 95
 Ser Tyr Pro Cys Lys Leu Leu Gln Met Ile Xaa Lys Ile Trp Xaa Pro
 100 105 110
 Cys Gly Xaa Leu Leu Thr Gly Gln Gln Xaa Ser Asn Glu Ile Ser Lys
 115 120 125
 Gln Glu Ile Xaa Cys Leu Leu His Pro Pro Pro Lys Asn Leu His Ile
 130 135 140
 Asp Val
 145

 <210> 159
 <211> 143
 <212> PRT
 <213> Homo sapiens

 <220>
 <221> SITE
 <222> (143)

<223> Xaa equals stop translation

<400> 159

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Met Trp Trp Ala Val Met Gly Gly Val Ile Gly Ser Trp Leu Ser Pro
 1           5           10           15

Leu Ser Ile Ala Glu Cys Cys His Asp Leu Trp Thr Ser Gln Ser Cys
 20           25           30

Glu His Ala Gly Ala Leu Cys Gly Asp Leu Leu Cys Ala Cys Arg Lys
 35           40           45

Val Gly Val Trp Cys Ala Leu Gln Gln His Trp Trp Asn Arg Cys Val
 50           55           60

Cys Pro His Ala Val Ile Arg Val His Cys Thr Gly Ala Ser Tyr Thr
 65           70           75           80

Leu Gln Lys Ile Cys Ser Cys Asn Pro Lys Phe Met Gly Arg His Pro
 85           90           95

His Arg Trp Gln Gln Ile Arg Lys Cys Ser Gln Pro Val Leu Arg Gly
100           105           110

Ser Arg Ala Ala Phe Ile Trp Val Arg Leu Ala Ala Leu Asn Phe Ile
115           120           125

Ser Ser Phe Arg Cys Ile Ser Leu Ile Ser Tyr Ser Ala Phe Xaa
130           135           140

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<210> 160

<211> 51

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (51)

<223> Xaa equals stop translation

<400> 160

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Met Lys Val Ser Asp Phe Asn Phe Leu Ile Phe Leu Ile Phe Ala Leu
 1           5           10           15

Phe Leu Thr Leu Glu Ala Phe Leu Lys Phe Thr Lys Arg Val Leu Ala
 20           25           30

Val Val Gly Asn Leu Pro Glu Pro Pro Ile Ile Lys Thr Ile Gly Phe
 35           40           45

Leu Tyr Xaa
 50

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<210> 161

<211> 65

<212> PRT

<213> Homo sapiens

<220>
<221> SITE
<222> (65)
<223> Xaa equals stop translation

<400> 161
Met Val Trp Ser Ala Ala Pro Ala Pro Cys Cys Leu Leu Gly Val Leu
1 5 10 15
Gly Leu Val Gln Val Leu Gly Ala Gln Ala Val Gly Pro Trp Thr Ala
20 25 30
Ser Ala Cys Leu Gly Ala Ala Gln Ala Gln Pro Cys Arg Pro Cys Lys
35 40 45
Glu Ser Ser Leu Arg Leu Phe Ser Ala Ser Ala Pro Ser Met Thr His
50 55 60
Xaa
65

<210> 162
<211> 59
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (59)
<223> Xaa equals stop translation

<400> 162
Met Glu Lys Tyr Cys Leu Gly Asn Asn Met Leu Ser Arg Phe Cys Leu
1 5 10 15
Phe Leu Ile Met Leu Leu His Ile Leu Leu Phe Leu Val Ile Phe Ile
20 25 30
Gln Arg His Thr Val Val Ser Leu Ser Lys His His Pro Phe Val Pro
35 40 45
Thr Asn Gly Ser Lys Ser Tyr Ser Ser Phe Xaa
50 55

<210> 163
<211> 374
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (84)
<223> Xaa equals any of the naturally occurring L-amino acids

<220>
<221> SITE

<222> (112)

<223> Xaa equals any of the naturally occurring L-amino acids

<400> 163

Met Arg Pro Gly Thr Ala Leu Gln Ala Val Leu Leu Ala Val Leu Leu
 1 5 10 15
 Val Gly Leu Arg Ala Ala Thr Gly Arg Leu Leu Ser Gly Gln Pro Val
 20 25 30
 Cys Arg Gly Gly Thr Gln Arg Pro Cys Tyr Lys Val Ile Tyr Phe His
 35 40 45
 Asp Thr Ser Arg Arg Leu Asn Phe Glu Glu Ala Lys Glu Ala Cys Arg
 50 55 60
 Arg Asp Gly Gly Gln Leu/Val Ser Ile Glu Ser Glu Asp Glu Gln Lys
 65 70 75 80
 Leu Ile Glu Xaa Phe Ile Glu Asn Leu Leu Pro Ser Asp Gly Asp Phe
 85 90 95
 Trp Ile Gly Leu Arg Arg Arg Glu Glu Lys Gln Ser Asn Ser Thr Xaa
 100 105 110
 Cys Gln Asp Leu Tyr Ala Trp Thr Asp Gly Ser Ile Ser Gln Phe Arg
 115 120 125
 Asn Trp Tyr Val Asp Glu Pro Ser Cys Gly Ser Glu Val Cys Val Val
 130 135 140
 Met Tyr His Gln Pro Ser Ala Pro Ala Gly Ile Gly Gly Pro Tyr Met
 145 150 155 160
 Phe Gln Trp Asn Asp Arg Cys Asn Met Lys Asn Asn Phe Ile Cys
 165 170 175
 Lys Tyr Ser Asp Glu Lys Pro Ala Val Pro Ser Arg Glu Ala Glu Gly
 180 185 190
 Glu Glu Thr Glu Leu Thr Thr Pro Val Leu Pro Glu Glu Thr Gln Glu
 195 200 205
 Glu Asp Ala Lys Lys Thr Phe Lys Glu Ser Arg Glu Ala Ala Leu Asn
 210 215 220
 Leu Ala Tyr Ile Leu Ile Pro Ser Ile Pro Leu Leu Leu Leu Val
 225 230 235 240
 Val Thr Thr Val Val Cys Trp Val Trp Ile Cys Arg Lys Arg Lys Arg
 245 250 255
 Glu Gln Pro Asp Pro Ser Thr Lys Lys Gln His Thr Ile Trp Pro Ser
 260 265 270
 Pro His Gln Gly Asn Ser Pro Asp Leu Glu Val Tyr Asn Val Ile Arg
 275 280 285
 Lys Gln Ser Glu Ala Asp Leu Ala Glu Thr Arg Pro Asp Leu Lys Asn

290 295 300
 Ile Ser Phe Arg Val Cys Ser Gly Glu Ala Thr Pro Asp Asp Met Ser
 305 310 315 320
 Cys Asp Tyr Asp Asn Met Ala Val Asn Pro Ser Glu Ser Gly Phe Val
 325 330 335
 Thr Leu Val Ser Val Glu Ser Gly Phe Val Thr Asn Asp Ile Tyr Glu
 340 345 350
 Phe Ser Pro Asp Gln Met Gly Arg Ser Lys Glu Ser Gly Trp Val Glu
 355 360 365
 Asn Glu Ile Tyr Gly Tyr
 370

<210> 164
 <211> 64
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (64)
 <223> Xaa equals stop translation

<400> 164
 Met His Pro Gln Leu Ile Pro Ser Val Ile Ala Val Val Phe Ile Leu
 1 5 10 15
 Leu Leu Gly Val Cys Phe Ile Ala Ser Cys Leu Val Thr His His Asn
 20 25 30
 Phe Ser Arg Cys Lys Arg Gly Thr Gly Val His Lys Leu Glu His His
 35 40 45
 Ala Lys Leu Lys Cys Ile Lys Glu Lys Ser Glu Leu Lys Ser Cys Xaa
 50 55 60

<210> 165
 <211> 743
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (743)
 <223> Xaa equals stop translation

<400> 165
 Met Ala Val Arg Glu Leu Cys Phe Pro Arg Gln Arg Gln Val Leu Phe
 1 5 10 15

Leu Phe Leu Phe Trp Gly Val Ser Leu Ala Gly Ser Gly Phe Gly Arg
 20 25 30
 Tyr Ser Val Thr Glu Glu Thr Glu Lys Gly Ser Phe Val Val Asn Leu
 35 40 45
 Ala Lys Asp Leu Gly Leu Ala Glu Gly Glu Leu Ala Ala Arg Gly Thr
 50 55 60
 Arg Val Val Ser Asp Asp Asn Lys Gln Tyr Leu Leu Leu Asp Ser His
 65 70 75 80
 Thr Gly Asn Leu Leu Thr Asn Glu Lys Leu Asp Arg Glu Lys Leu Cys
 85 90 95
 Gly Pro Lys Glu Pro Cys Met Leu Tyr Phe Gln Ile Leu Met Asp Asp
 100 105 110
 Pro Phe Gln Ile Tyr Arg Ala Glu Leu Arg Val Arg Asp Ile Asn Asp
 115 120 125
 His Ala Pro Val Phe Gln Asp Lys Glu Thr Val Leu Lys Ile Ser Glu
 130 135 140
 Asn Thr Ala Glu Gly Thr Ala Phe Arg Leu Glu Arg Ala Gln Asp Pro
 145 150 155 160
 Asp Gly Gly Leu Asn Gly Ile Gln Asn Tyr Thr Ile Ser Pro Asn Ser
 165 170 175
 Phe Phe His Ile Asn Ile Ser Gly Gly Asp Glu Gly Met Ile Tyr Pro
 180 185 190
 Glu Leu Val Leu Asp Lys Ala Leu Asp Arg Glu Glu Gln Gly Glu Leu
 195 200 205
 Ser Leu Thr Leu Thr Ala Leu Asp Gly Gly Ser Pro Ser Arg Ser Gly
 210 215 220
 Thr Ser Thr Val Arg Ile Val Val Leu Asp Val Asn Asp Asn Ala Pro
 225 230 235 240
 Gln Phe Ala Gln Ala Leu Tyr Glu Thr Gln Ala Pro Glu Asn Ser Pro
 245 250 255
 Ile Gly Phe Leu Ile Val Lys Val Trp Ala Glu Asp Val Asp Ser Gly
 260 265 270
 Val Asn Ala Glu Val Ser Tyr Ser Phe Phe Asp Ala Ser Glu Asn Ile
 275 280 285
 Arg Thr Thr Phe Gln Ile Asn Pro Phe Ser Gly Glu Ile Phe Leu Arg
 290 295 300
 Glu Leu Leu Asp Tyr Glu Leu Val Asn Ser Tyr Lys Ile Asn Ile Gln
 305 310 315 320
 Ala Met Asp Gly Gly Gly Leu Ser Ala Arg Cys Arg Val Leu Val Glu
 325 330 335

Val Leu Asp Thr Asn Asp Asn Pro Pro Glu Leu Ile Val Ser Ser Phe
340 345 350
Ser Asn Ser Val Ala Glu Asn Ser Pro Glu Thr Pro Leu Ala Val Phe
355 360 365
Lys Ile Asn Asp Arg Asp Ser Gly Glu Asn Gly Lys Met Val Cys Tyr
370 375 380
Ile Gln Glu Asn Leu Pro Phe Leu Leu Lys Pro Ser Val Glu Asn Phe
385 390 395 400
Tyr Ile Leu Ile Thr Glu Gly Ala Leu Asp Arg Glu Ile Arg Ala Glu
405 410 415
Tyr Asn Ile Thr Ile Thr Val Thr Asp Leu Gly Thr Pro Arg Leu Lys
420 425 430
Thr Glu His Asn Ile Thr Val Leu Val Ser Asp Val Asn Asn Asn Ala
435 440 445
Pro Ala Phe Thr Gln Thr Ser Tyr Thr Leu Phe Val Arg Glu Asn Asn
450 455 460
Ser Pro Ala Leu His Ile Gly Ser Val Ser Ala Thr Asp Arg Asp Ser
465 470 475 480
Gly Thr Asn Ala Gln Val Thr Tyr Ser Leu Leu Pro Pro Gln Asp Pro
485 490 495
His Leu Pro Leu Ala Ser Leu Val Ser Ile Asn Ala Asp Asn Gly His
500 505 510
Leu Phe Ala Leu Arg Ser Leu Asp Tyr Glu Ala Leu Gln Ala Phe Glu
515 520 525
Phe Arg Val Gly Ala Thr Asp Arg Gly Ser Pro Ala Leu Asn Ser Glu
530 535 540
Ala Leu Gly Ala Arg Ala Gly Ala Gly Arg Gln Arg Gln Leu Ala Leu
545 550 555 560
Arg Ala Val Pro Ala Ala Glu Arg Leu Arg Ala Leu His Arg Ala Gly
565 570 575
Ala Pro Gly Gly Arg Ala Gly Leu Pro Gly Asp Gln Gly Gly Gly Gly
580 585 590
Gly Arg Arg Leu Gly Pro Glu Arg Leu Ala Val Val Pro Ala Ala Gln
595 600 605
Gly His Gly Ala Arg Ala Val Arg Cys Val Gly Ala Gln Trp Gly Gly
610 615 620
Ala His Arg Gln Ala Ala Glu Arg Ala Arg Arg Ser Gln Ala Gln Ala
625 630 635 640
Gly Gly Ala Cys Gln Gly Gln Trp Arg Ala Ser Ser Leu Gly His Arg

645 650 655
 His Ala Ala Arg Ala Pro Gly Gly Arg Leu Leu Pro Ala Leu Pro Ala
 660 665 670
 Ser Pro Gly Gly Gly Pro Gly Pro Gly Pro Gly Arg Leu Ala His Arg
 675 680 685
 Leu Pro Gly Gly Gly Val Gly Leu Gly Val Phe Ala Leu Pro Pro Leu
 690 695 700
 Gly Ala Pro Val Arg Gly Gly Ala Ala Val Gln Glu Glu Gln Gly Gly
 705 710 715 720
 Leu Gly Gly Ser Leu Leu Gly Ala Arg Gly Ser Phe Ser Arg Ala Ser
 725 730 735
 Gly Gly Arg Glu Gly Arg Xaa
 740

<210> 166
 <211> 214
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (214)
 <223> Xaa equals stop translation

<400> 166
 Met Asn Arg Met Glu Leu Leu Lys Leu Leu Leu Thr Cys Phe Ser Glu
 1 5 10 15
 Ala Met Tyr Leu Pro Pro Ala Pro Glu Ser Gly Ser Thr Asn Pro Trp
 20 25 30
 Val Gln Phe Phe Cys Ser Thr Glu Asn Arg His Ala Leu Pro Leu Phe
 35 40 45
 Thr Ser Leu Leu Asn Thr Val Cys Ala Tyr Asp Pro Val Gly Tyr Gly
 50 55 60
 Ile Pro Tyr Asn His Leu Leu Phe Ser Asp Tyr Arg Glu Pro Leu Val
 65 70 75 80
 Glu/Glu Ala Ala Gln Val Leu Ile Val Thr Leu Asp His Asp Ser Ala
 85 90 95
 Ser Ser Ala Ser Pro Thr Val Asp Gly Thr Thr Thr Gly Thr Ala Met
 100 105 110
 Asp Asp Ala Asp Pro Pro Gly Pro Glu Asn Leu Phe Val Asn Tyr Leu
 115 120 125
 Ser Arg Ile His Arg Glu Glu Asp Phe Gln Phe Ile Leu Lys Gly Ile
 130 135 140

Ala Arg Leu Leu Ser Asn Pro Leu Leu Gln Thr Tyr Leu Pro Asn Ser
 145 150 155 160
 Thr Lys Lys Asp Pro Val Pro Pro Gly Ala Ala Ser Ser Leu Leu Glu
 165 170 175
 Ala Leu Arg Leu Gln Gln Glu Ile Pro Leu Leu Arg Ala Glu Glu Gln
 180 185 190
 Arg Arg Pro Arg His Pro Cys Pro His Pro Leu Leu Pro Gln Arg Cys
 195 200 205
 Pro Gly Arg Ser Val Xaa
 210

<210> 167
 <211> 213
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (213)
 <223> Xaa equals stop translation

<400> 167
 Met Pro Ser Leu Arg Phe Leu Ala Leu Ala Leu Leu Ala Ile Leu
 1 5 10 15
 Pro Ala Leu Pro Asn Ala His Ala Ala Pro Gly Ile Gly Gly Leu Ile
 20 25 30
 Gly Gly Gly Ser Gln Ala Ser Ala Lys Glu Glu Pro Gln Ser Asn Ala
 35 40 45
 Gln Pro Ser Ala Asp Glu Arg Lys Gln Arg Leu Leu Ser Gln Ala Glu
 50 55 60
 Glu Thr Arg Gln Arg Leu Thr Asp Leu Lys Ala Glu Leu Ala Gly Ala
 65 70 75 80
 Pro Lys Glu Ile Ser Glu Ala Gln Arg Thr Leu Ser Lys Leu Val Ser
 85 90 95
 Glu Asp Asn Ser Asp Leu Pro Glu Arg Leu Ser Lys Leu Ser Val Pro
 100 105 110
 Val Leu Glu Gln Arg Leu Ala Ala Arg Val Asp Glu Leu Ala Leu Trp
 115 120 125
 Gln Gln Ala Leu Ser Ala Ala Asn Ser Met Leu Ile Ser Ala Gln Thr
 130 135 140
 Arg Pro Glu Arg Ala Gln Ala Asp Ile Ser Lys Asn Gln Leu Arg Ile
 145 150 155 160
 Asp Glu Ile Asn Gly Leu Leu Lys Ser Gly Arg Glu Asn Asn Lys Pro
 165 170 175

Leu Thr Asp Glu Arg Arg Ala Leu Leu Glu Ser Thr Ser Arg Ala Ala
180 185 190

Ala Gly Pro Ser Ile Phe His Pro Gly Gly Val Pro Gly Lys Cys Thr
195 200 205

Gln Phe Ala Leu Xaa
210

<210> 168

<211> 75

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (75)

<223> Xaa equals stop translation

<400> 168

Met Phe Thr Ser Phe Gly Leu Ala Ser Pro Arg Ile Leu Phe Cys Phe
1 5 10 15

Cys Phe Phe Asp Leu Gly Phe Ile Phe Phe Cys Val Leu Tyr Tyr Ile
20 25 30

Val Lys Gly Ile Leu Ala Glu Thr Leu Val Phe Gly Ala Arg Gly Glu
35 40 45

Gln Glu Cys Trp Ala Val Tyr Phe Arg Trp Arg Thr His Leu Gln Thr
50 55 60

Phe Gly Leu Phe Ser Phe Asn Cys Ser Val Xaa
65 70 75

<210> 169

<211> 48

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (48)

<223> Xaa equals stop translation

<400> 169

Met Phe Leu Cys Leu Phe Phe Phe Phe Phe Asn Ala Thr Gln Gly Asn
1 5 10 15

Ile Phe Ile Ser Phe Leu Ser Gly Leu Pro Gln Cys Ile Phe Ile Ser
20 25 30

Phe Glu Thr Lys Arg Phe Trp Lys Leu Phe Phe Cys Ser Phe Lys Xaa
35 40 45

<210> 170
<211> 88
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (88)
<223> Xaa equals stop translation

<400> 170
Met Gly Leu His Leu Arg Pro Tyr Arg Val Gly Leu Leu Pro Asp Gly
1 5 10 15
Leu Leu Phe Leu Leu Leu Leu Met Leu Leu Ala Asp Pro Ala Leu
20 25 30
Pro Ala Gly Arg His Pro Pro Val Val Leu Val Pro Gly Asp Leu Gly
35 40 45
Asn Gln Leu Glu Ala Lys Leu Asp Lys Pro Thr Val Val His Tyr Leu
50 55 60
Cys Ser Lys Lys Thr Glu Ser Tyr Phe Thr Ile Trp Leu Asn Leu Glu
65 70 75 80
Leu Leu Leu Pro Val His His Xaa
85

<210> 171
<211> 42
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (42)
<223> Xaa equals stop translation

<400> 171
Met Ala Cys Glu Thr His Gly Val Leu Val Pro Ala His Leu Ser Gly
1 5 10 15
Leu Ile Thr Cys Leu Leu Ala Phe Trp Val Pro Ala Ser Cys Ile Gln
20 25 30
Arg Cys Ser Gly Ser Pro Leu Pro Leu Xaa
35 40

<210> 172
<211> 48
<212> PRT
<213> Homo sapiens

<220>
 <221> SITE
 <222> (48)
 <223> Xaa equals stop translation

<400> 172
 Met Gln Cys Phe Leu Phe Ser Ile Phe Leu Ile Thr Gly Leu Ala Glu
 1 5 10 15
 Glu Phe Cys Glu Gln Leu Ser Ile Ser Leu Ala Glu Glu Glu Ile Gln
 20 25 30
 Leu Ser Ser Thr Val Glu His Phe Cys Met Thr Ala Phe Ser Trp Xaa
 35 40 45

<210> 173
 <211> 233
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (233)
 <223> Xaa equals stop translation

<400> 173
 Met Ala Ala Leu Ala Ala Ala Lys Lys Val Trp Ser Ala Arg Arg
 1 5 10 15
 Leu Leu Val Leu Leu Phe Thr Pro Leu Ala Leu Leu Pro Val Val Phe
 20 25 30
 Ala Leu Pro Pro Lys Glu Gly Arg Cys Leu Phe Val Ile Leu Leu Met
 35 40 45
 Ala Val Tyr Trp Cys Thr Glu Ala Leu Pro Leu Ser Val Thr Ala Leu
 50 55 60
 Leu Pro Ile Val Leu Phe Pro Phe Met Gly Ile Leu Pro Ser Asn Lys
 65 70 75 80
 Val Cys Pro Gln Tyr Phe Leu Asp Thr Asn Phe Leu Phe Leu Ser Gly
 85 90 95
 Leu Ile Met Ala Ser Ala Ile Glu Glu Trp Asn Leu His Arg Arg Ile
 100 105 110
 Ala Leu Lys Ile Leu Met Leu Val Gly Val Gln Pro Ala Arg Leu Ile
 115 120 125
 Leu Gly Met Met Val Thr Thr Ser Phe Leu Ser Met Trp Leu Ser Asn
 130 135 140
 Thr Ala Ser Thr Ala Met Met Leu Pro Ile Ala Asn Ala Ile Leu Lys
 145 150 155 160

Ser Leu Phe Gly Gln Lys Glu Val Arg Lys Asp Pro Ser Gln Glu Ser
 165 170 175
 Glu Glu Asn Thr Gly Ile Glu Pro Asn Thr Phe Leu Ser Glu Glu Arg
 180 185 190
 Leu Lys Leu Gln Ala Pro Leu Val Ile Arg Leu Gly Gln Ile Thr Glu
 195 200 205
 Ser Gly Gln Trp Asn Met Ser Gly Asn Asp Val Cys Asn Phe Arg Val
 210 215 220
 Leu Ser Phe Leu Pro Gly Gly Met Xaa
 225 230

<210> 174
 <211> 45
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (45)
 <223> Xaa equals stop translation

<400> 174
 Met Gly Thr Ile Phe Gly Tyr Leu His Cys Val Lys Cys Tyr Val Leu
 1 5 10 15
 Tyr Phe Ile Phe Ile Leu Ile Thr Ala Val Tyr His Ser Phe Tyr Tyr
 20 25 30
 Pro His Tyr Arg Gly Lys Ala Leu Ile Ser Gly Thr Xaa
 35 40 45

<210> 175
 <211> 85
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (77)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (85)
 <223> Xaa equals stop translation

<400> 175
 Met Val Trp Phe Leu Phe Leu Val Phe Ile Phe Leu Lys Val Lys Gly
 1 5 10 15
 Asp Phe Phe Pro Pro Phe Leu Ile Cys Asn Leu Phe Cys Ile Trp Met
 20 25 30

Ile Thr Gly Val Ser His Arg Leu Gln Pro Gln Ile Leu Phe Ser Arg
 35 40 45
 His Lys His Asn Gln Glu Ile Ile Leu Gln Met Val Ser Phe Ser Cys
 50 55 60
 Cys Val Phe Phe Pro Met Ile Arg Glu Val Lys Ser Xaa Leu Gly Cys
 65 70 75 80
 Ile Lys Met Ser Xaa
 85

<210> 176
 <211> 66
 <212> PRT
 <213> Homo sapiens
 <220>
 <221> SITE
 <222> (66)
 <223> Xaa equals stop translation

<400> 176
 Met Trp Val Leu Leu Ser Cys Pro Leu Pro Pro Leu Cys Leu Pro Ala
 1 5 10 15
 Ser Ala Val Pro Gly Gln Cys Leu Gly Gly Gln Trp Ser Gly His Gln
 20 25 30
 Leu Arg Leu Arg Gly Arg Gly Trp His Cys Arg Cys His Cys Arg Ala
 35 40 45
 Trp Ala Ala Asp Met Gly Arg Gly Leu His Ser Cys Gln Leu Leu Ser
 50 55 60
 Arg Xaa
 65

<210> 177
 <211> 55
 <212> PRT
 <213> Homo sapiens
 <220>
 <221> SITE
 <222> (55)
 <223> Xaa equals stop translation

<400> 177
 Met Leu Leu Leu Cys Ile Leu Leu Ile Phe Cys Val Val Gly Leu Ser
 1 5 10 15
 Val Val Gly Arg Arg Val Leu Lys Ser Thr Thr Ile Ile Val Tyr Leu
 20 25 30
 Ser Ile Thr Pro Phe Ser Ser Phe Ser Ser Ile Ser His Ile Phe Gln

35 40 45

Leu Leu Ile Gly Ala His Xaa
50 55

<210> 178
<211> 83
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (4)
<223> Xaa equals any of the naturally occurring L-amino acids

<220>
<221> SITE
<222> (83)
<223> Xaa equals stop translation

<400> 178
Met Cys Val Xaa Leu Ser Phe Cys Pro Phe Leu Ser Ser Ala Leu Pro
1 5 10 15
Ala Ser His Thr Gln Phe Tyr Met Pro Arg Gly Ala Lys Phe Gly Thr
20 25 30
Phe Thr Leu Gln Ala Ser Val Ser Pro Leu Glu Glu Lys Thr His Ser
35 40 45
Phe Thr His Pro Gly Ile Gly Gly Lys Leu Leu Gly His Gln Asp Pro
50 55 60
Gly Ala Pro Gly Pro Ser Trp Asn Ile Arg Ser Thr Trp Ser Thr Arg
65 70 75 80
Ser Leu Xaa

<210> 179
<211> 330
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (38)
<223> Xaa equals any of the naturally occurring L-amino acids

<220>
<221> SITE
<222> (247)
<223> Xaa equals any of the naturally occurring L-amino acids

<400> 179
Met Ser Pro Leu Ser Ala Ala Arg Ala Ala Leu Arg Val Tyr Ala Val
1 5 10 15

Gly Ala Ala Val Ile Leu Ala Gln Leu Leu Arg Arg Cys Arg Gly Gly
 20 25 30
 Phe Leu Glu Pro Val Xaa Pro Pro Arg Pro Asp Arg Val Ala Ile Val
 35 40 45
 Thr Gly Gly Thr Asp Gly Ile Gly Tyr Ser Thr Ala Lys His Leu Ala
 50 55 60
 Arg Leu Gly Met His Val Ile Ile Ala Gly Asn Asn Asp Ser Lys Ala
 65 70 75 80
 Lys Gln Val Val Ser Lys Ile Lys Glu Glu Thr Leu Asn Asp Lys Val
 85 90 95
 Glu Phe Leu Tyr Cys Asp Leu Ala Ser Met Thr Ser Ile Arg Gln Phe
 100 105 110
 Val Gln Lys Phe Lys Met Lys Lys Ile Pro Leu His Val Leu Ile Asn
 115 120 125
 Asn Ala Gly Val Met Met Val Pro Gln Arg Lys Thr Arg Asp Gly Phe
 130 135 140
 Glu Glu His Phe Gly Leu Asn Tyr Leu Gly His Phe Leu Leu Thr Asn
 145 150 155 160
 Leu Leu Leu Asp Thr Leu Lys Glu Ser Gly Ser Pro Gly His Ser Ala
 165 170 175
 Arg Val Val Thr Val Ser Ser Ala Thr His Tyr Val Ala Glu Leu Asn
 180 185 190
 Met Asp Asp Leu Gln Ser Ser Ala Cys Tyr Ser Pro His Ala Ala Tyr
 195 200 205
 Ala Gln Ser Lys Leu Ala Leu Val Leu Phe Thr Tyr His Leu Gln Arg
 210 215 220
 Leu Leu Ala Ala Glu Gly Ser His Val Thr Ala Asn Val Val Asp Pro
 225 230 235 240
 Gly Val Val Asn Thr Asp Xaa Tyr Lys His Val Phe Trp Ala Thr Arg
 245 250 255
 Leu Ala Lys Lys Leu Leu Gly Trp Leu Leu Phe Lys Thr Pro Asp Glu
 260 265 270
 Gly Ala Trp Thr Ser Ile Tyr Ala Ala Val Thr Pro Glu Leu Glu Gly
 275 280 285
 Val Gly Gly Arg Tyr Leu Tyr Asn Gln Lys Glu Thr Lys Ser Leu His
 290 295 300
 Val Thr Tyr Asn Gln Lys Leu Gln Gln Gln Leu Trp Ser Lys Ser Cys
 305 310 315 320
 Glu Met Thr Gly Val Leu Asp Val Thr Leu

325

330

<210> 180
<211> 41
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (41)
<223> Xaa equals stop translation

<400> 180
Met Ile Ala Cys Gln Tyr Ile Ser Leu Ala Ile Met Leu Ala Phe Val
1 5 10 15
Arg Trp Ala Ala Phe Leu Leu Phe Pro Phe Leu Cys Gly Asp Asn Gly
20 25 30
Gly Asn Ile Gln Gln Lys Tyr Val Xaa
35 40

<210> 181
<211> 52
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (52)
<223> Xaa equals stop translation

<400> 181
Met Ala Asn Ala Met Ala Tyr Leu Ser Ile Phe Leu Cys Gly Ala Ser
1 5 10 15
Ser Ser Pro Cys Asp Cys Ala Leu Leu Val Pro Val Ser Leu Phe Arg
20 25 30
Gly Arg Lys Val Ala Asn Phe Lys Asn Gln Asn Ser Asp Val Thr Ser
35 40 45
Gly Asn Ala Xaa
50

<210> 182
<211> 55
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (55)
<223> Xaa equals stop translation

<400> 182

Met Gln Gln Ile Cys Ser Cys Leu Gly Ala Phe Ala Leu Leu Phe Phe
 1 5 10 15
 Trp Pro Gly His Phe Thr Ser Thr Phe Ser Ile Phe Tyr Asp Phe Leu
 20 25 30
 Pro Ile Phe Gly Ser Leu Phe Lys Cys His Pro Ser Lys Arg Pro Ser
 35 40 45
 Lys Leu Pro Tyr Leu Lys Xaa
 50 55

<210> 183
 <211> 62
 <212> PRT
 <213> Homo sapiens
 <220>
 <221> SITE
 <222> (62)
 <223> Xaa equals stop translation

<400> 183
 Met Arg Leu Leu Leu Glu Trp Arg Val Tyr Leu Arg Leu Thr Cys Ala
 1 5 10 15
 Thr Lys Asp Gly Met Ala Arg Glu Cys Pro Thr Thr Trp Leu Ser Pro
 20 25 30
 Pro Ala Lys Pro Asp Phe Ala Gln Arg His Ser Val Lys Pro Thr Ala
 35 40 45
 Leu Gln Gly Gly Arg Trp Ser Arg Leu Gly Ala Ser Pro Xaa
 50 55 60

<210> 184
 <211> 148
 <212> PRT
 <213> Homo sapiens
 <220>
 <221> SITE
 <222> (148)
 <223> Xaa equals stop translation

<400> 184
 Met Leu Gly Leu Pro Trp Lys Gly Gly Leu Ser Trp Ala Leu Leu Leu
 1 5 10 15
 Leu Leu Leu Gly Ser Gln Ile Leu Leu Ile Tyr Ala Trp His Phe His
 20 25 30
 Glu Gln Arg Asp Cys Asp Glu His Asn Val Met Ala Arg Tyr Leu Pro
 35 40 45
 Ala Thr Val Glu Phe Ala Val His Thr Phe Asn Gln Gln Ser Lys Asp
 50 55 60

Tyr Tyr Ala Tyr Arg Leu Gly His Ile Leu Asn Ser Trp Lys Glu Gln
 65 70 75 80
 Val Glu Ser Lys Thr Val Phe Ser Met Glu Leu Leu Leu Gly Arg Thr
 85 90 95
 Arg Cys Gly Lys Phe Glu Asp Asp Ile Asp Asn Cys His Phe Gln Glu
 100 105 110
 Ser Thr Glu Leu Asn Asn Thr Phe Thr Cys Phe Phe Thr Ile Ser Thr
 115 120 125
 Arg Pro Trp Met Thr Gln Phe Ser Leu Leu Asn Lys Thr Cys Leu Glu
 130 135 140
 Gly Phe His Xaa
 145

<210> 185
 <211> 161
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (146)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (151)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (161)
 <223> Xaa equals stop translation

<400> 185
 Met Arg Leu Leu Cys Gly Leu Trp Leu Trp Leu Ser Leu Leu Lys Val
 1 5 10 15
 Leu Gln Ala Gln Thr Pro Thr Pro Leu Pro Leu Pro Pro Pro Met Gln
 20 25 30
 Ser Phe Gln Gly Asn Gln Phe Gln Gly Glu Trp Phe Val Leu Gly Leu
 35 40 45
 Ala Gly Asn Ser Phe Arg Pro Glu His Arg Ala Leu Leu Asn Ala Phe
 50 55 60
 Thr Ala Thr Phe Glu Leu Ser Asp Asp Gly Arg Phe Glu Val Trp Asn
 65 70 75 80
 Ala Met Thr Arg Gly Gln His Cys Asp Thr Trp Ser Tyr Val Leu Ile
 85 90 95

Pro Ala Ala Gln Pro Gly Gln Phe Thr Val Asp His Gly Val Gly Arg
100 105 110
Ser Trp Leu Leu Pro Pro Gly Thr Leu Asp Gln Phe Ile Cys Leu Gly
115 120 125
Arg Ala Gln Gly Leu Ser Asp Asp Asn Ile Val Phe Pro Asp Val Thr
130 135 140
Gly Xaa Ala Leu Asp Leu Xaa Ser Leu Pro Trp Val Ala Ala Pro Ala
145 150 155 160
Xaa

<210> 186
<211> 122
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (122)
<223> Xaa equals stop translation

<400> 186
Met Met Leu Pro Gln Trp Leu Leu Leu Phe Leu Leu Phe Phe Phe
1 5 10 15
Leu Phe Leu Leu Thr Arg Gly Ser Leu Ser Pro Thr Lys Tyr Asn Leu
20 25 30
Leu Glu Leu Lys Glu Ser Cys Ile Arg Asn Glr Asp Cys Glu Thr Gly
35 40 45
Cys Cys Gln Arg Ala Pro Asp Asn Cys Glu Ser His Cys Ala Glu Lys
50 55 60
Gly Ser Glu Gly Ser Leu Cys Gln Thr Gln Val Phe Phe Gly Gln Tyr
65 70 75 80
Arg Ala Cys Pro Cys Leu Arg Asn Leu Thr Cys Ile Tyr Ser Lys Asn
85 90 95
Glu Lys Trp Leu Ser Ile Ala Tyr Gly Arg Cys Gln Lys Ile Gly Arg
100 105 110
Gln Lys Leu Ala Lys Lys Met Phe Phe Xaa
115 120

<210> 187
<211> 163
<212> PRT
<213> Homo sapiens

<220>
<221> SITE

<222> (163)

<223> Xaa equals stop translation

<400> 187

Met Thr Ser Asn Phe Pro Phe Cys Thr Leu Ile Leu Gly Ile Ala Gln
 1 5 10 15

Ala Gln Ala Cys Pro Gly Cys Pro Gly Asp Trp Pro Gly Leu Gly Ser
 20 25 30

Gly Val Gly Glu Gly Leu His His Ile Arg Thr Cys Arg Thr Pro Ile
 35 40 45

Pro Cys Ser Pro Pro Ala Pro Ala Ala Ala Cys Leu Gly Ser Gly His
 50 55 60

Ala Arg Leu Pro Cys Val Leu Arg Leu Trp Pro Val Pro Ala Asn Leu
 65 70 75 80

Ser Ser Pro Phe Arg Leu Glu Ala Leu His Cys Ser Phe Trp Ser Ser
 85 90 95

Pro Leu Leu Pro Ala Pro His Leu Ala Phe Phe Gly Phe Arg Asp Leu
 100 105 110

Leu Thr Asp Phe Leu Leu Ala Ala Cys Leu Leu Thr Phe Gln Lys Thr
 115 120 125

Pro Leu Glu Leu Pro Met Ala Val Val His Leu Leu Val Ala Thr Pro
 130 135 140

Cys Tyr Gln Met Leu Asp Asn Leu Pro Leu Pro Ser Ala Ala Ala Asn
 145 150 155 160

Trp Cys Xaa

<210> 188

<211> 51

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (51)

<223> Xaa equals stop translation

<400> 188

Met Pro Gly Ile Leu Ala Gly Ile Pro Val Lys Asp Leu Cys Leu Ser
 1 5 10 15

Leu Leu Gln Gly Phe Arg Leu Leu Leu Leu Cys Val Cys Pro Gly Trp
 20 25 30

Leu Ser Gly Trp Met Gly Gly Gln Lys Gly Ser Pro Arg Ile Val Asp
 35 40 45

Ile Gly Xaa

50

<210> 189
<211> 65
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (65)
<223> Xaa equals stop translation

<400> 189
Met Tyr Leu Tyr Leu Gly Val Phe Phe His Leu Ile Tyr Pro Gly Ala
1 5 10 15
Leu Ser Ile Thr Thr Leu Gly Lys His Ser His Pro Phe Thr Ala
20 25 30
Glu Gln Asn Ser Thr Val Trp Met Glu His Thr Leu Phe His Gln Ser
35 40 45
Pro Val Ala Ser His Leu Val Cys Phe Gln Ser Phe Ala Phe Ser Glu
50 55 60
Xaa
65

<210> 190
<211> 47
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (47)
<223> Xaa equals stop translation

<400> 190
Met Thr Leu Ser Leu Gln Leu Ala Glu Leu Val His Phe Val Cys Ala
1 5 10 15
Phe Gln Ser Gln Trp Thr Gly Val Tyr Pro Met Met Pro Pro Leu Lys
20 25 30
Pro Thr Glu Pro Leu Cys Phe Ala Cys Val Pro Cys Arg Val Xaa
35 40 45

<210> 191
<211> 144
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (144)

<223> Xaa equals stop translation

<400> 191

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Met Ser Pro Phe His Leu Leu Gly Leu Lys Val Phe Leu Thr Trp Ala
 1           5           10           15

Leu Thr Leu Ala Gln Ile Cys Leu Tyr Phe Phe Glu Val Gln Pro Leu
      20           25           30

Gly Leu Leu Ala Leu Asn Phe Phe Cys Thr Ala Thr Ala Gly Leu Lys
      35           40           45

Glu Leu Cys Met His Pro Pro Ser Leu Ala Phe Thr Pro Glu Phe His
      50           55           60

Thr Ser Leu Ser Pro Leu Ala Ile Pro Ser Phe Cys Gly Thr Ser Val
      65           70           75           80

Ser Leu Ser Asn Ser His Thr Ile Pro Leu Ser Leu Tyr Leu Pro Phe
      85           90           95

Pro Ser Lys Ser Arg Met Pro Asp Thr Leu His Leu Leu Val His Ser
      100          105          110

Leu Pro Leu Val His Ser Gln Val Leu Pro Val Lys Asp Val Thr Ile
      115          120          125

Glu Trp Pro Leu Cys Gln Arg Cys Leu Gly Ser Thr Cys His Gln Xaa
      130          135          140

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<210> 192

<211> 81

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (76)

<223> Xaa equals any of the naturally occurring L-amino acids

<220>

<221> SITE

<222> (81)

<223> Xaa equals any of the naturally occurring L-amino acids

<400> 192

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Met Phe Cys Phe Ser Ser Ile Phe Cys Ser His Glu His Thr His Leu
 1           5           10           15

Pro Gly Thr Phe Trp Leu Phe Leu Phe Leu Ile Leu Pro Pro
      20           25           30

Ser Cys Pro Cys Phe Leu Pro Phe Ser Leu Ala Ile Glu Thr Val Arg
      35           40           45

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Trp Pro Cys Trp His His Pro Thr Ser Phe Glu Leu Cys Tyr Pro Gly
50 55 60
Thr Ser Ile Tyr Tyr Ala Ser Arg Gly Gly Pro Xaa Pro Asn Ser Glu
65 70 75 80
Xaa

<210> 193
<211> 45
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (45)
<223> Xaa equals stop translation

<400> 193
Met Thr Tyr Leu Phe Cys Ser Ser Ile Ser Leu Leu Leu Lys Val
1 5 10 15
His Ser Ser Gly His Gln Asp Ile Arg Lys Ala Lys Ser Lys Val Pro
20 25 30
Arg Leu Leu Ile Ile Gln Cys Pro Gln Gln Arg Glu Xaa
35 40 45

<210> 194
<211> 42
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (42)
<223> Xaa equals stop translation

<400> 194
Met Pro Thr Ile Trp Val Lys Leu Cys Leu Leu Gln Val Cys His Gly
1 5 10 15
Leu Phe Pro Leu Leu Lys His Trp Ser Gln Pro Met Pro Leu Cys Val
20 25 30
Thr Leu Ala Pro Val Ser Tyr Trp Leu Xaa
35 40

<210> 195
<211> 260
<212> PRT
<213> Homo sapiens

<220>
<221> SITE

<222> (260)

<223> Xaa equals stop translation

<400> 195

Met Gly Thr Ala Ala Leu Gly Pro Val Trp Ala Ala Leu Leu Leu Phe
 1 5 10 15

Leu Leu Met Cys Glu Ile Pro Met Val Glu Leu Thr Phe Asp Arg Ala
 20 25 30

Val Ala Ser Asp Cys Gln Arg Cys Cys Asp Ser Glu Asp Pro Leu Asp
 35 40 45

Pro Ala His Val Ser Ser Ala Ser Ser Ser Gly Arg Pro His Ala Leu
 50 55 60

Pro Glu Ile Arg Pro Tyr Ile Asn Ile Thr Ile Leu Lys Gly Asp Lys
 65 70 75 80

Gly Asp Pro Gly Pro Met Gly Leu Pro Gly Tyr Met Gly Arg Glu Gly
 85 90 95

Pro Gln Gly Glu Pro Gly Pro Gln Gly Ser Lys Gly Asp Lys Gly Glu
 100 105 110

Met Gly Ser Pro Gly Ala Pro Cys Gln Lys Arg Phe Phe Ala Phe Ser
 115 120 125

Val Gly Arg Lys Thr Ala Leu His Ser Gly Glu Asp Phe Gln Thr Leu
 130 135 140

Leu Phe Glu Arg Val Phe Val Asn Leu Asp Gly Cys Phe Asp Met Ala
 145 150 155 160

Thr Gly Gln Phe Ala Ala Pro Leu Arg Gly Ile Tyr Phe Phe Ser Leu
 165 170 175

Asn Val His Ser Trp Asn Tyr Lys Glu Thr Tyr Val His Ile Met His
 180 185 190

Asn Gln Lys Glu Ala Val Ile Leu Tyr Ala Gln Pro Ser Glu Arg Ser
 195 200 205

Ile Met Gln Ser Gln Ser Val Met Leu Asp Leu Ala Tyr Gly Asp Arg
 210 215 220

Val Trp Val Arg Leu Phe Lys Arg Gln Arg Glu Asn Ala Ile Tyr Ser
 225 230 235 240

Asn Asp Phe Asp Thr Tyr Ile Thr Phe Ser Gly His Leu Ile Lys Ala
 245 250 255

Glu Asp Asp Xaa
 260

<210> 196

<211> 117

<212> PRT

<213> Homo sapiens

<400> 196

Met Leu Gly His Cys Cys Tyr Phe Trp Gln Val Trp Pro Ala Ser Glu
 1 5 10 15
 Ala Leu Ala Ala Gly Pro Thr Pro Ser Thr Gly Ser Ser Ser Pro Ser
 20 25 30
 Trp Lys Gln His Ile Gly Thr Ser Leu Gln Lys Thr Arg Gly Ser Leu
 35 40 45
 Pro Thr Thr Thr Leu Thr Ser Gly Ala Gly Gln Ser Thr Ser Thr Gly
 50 55 60
 Lys Asn Pro Ala Ala Gly Arg Ser Leu Glu Gly Ala Leu Pro Ala Gly
 65 70 75 80
 Val Trp Pro Cys Phe Ala Gln Ser Pro Cys Thr Gly Gly Gln Gln Thr
 85 90 95
 Pro Ser Ser Thr Gly Leu Arg Ser Cys Leu Val Arg Ser Pro Ala Thr
 100 105 110
 Trp Trp Arg Thr Pro
 115

<210> 197

<211> 698

<212> PRT

<213> Homo sapiens

<400> 197

Met Leu Pro Ala Arg Leu Pro Phe Arg Leu Leu Ser Leu Phe Leu Arg
 1 5 10 15
 Gly Ser Ala Pro Thr Ala Ala Arg His Gly Leu Arg Glu Pro Leu Leu
 20 25 30
 Glu Arg Arg Cys Ala Ala Ala Ser Ser Phe Gln His Ser Ser Ser Leu
 35 40 45
 Gly Arg Glu Leu Pro Tyr Asp Pro Val Asp Thr Glu Gly Phe Gly Glu
 50 55 60
 Gly Gly Asp Met Gln Glu Arg Phe Leu Phe Pro Glu Tyr Ile Leu Asp
 65 70 75 80
 Pro Glu Pro Gln Pro Thr Arg Glu Lys Gln Leu Gln Glu Leu Gln Gln
 85 90 95
 Gln Gln Glu Glu Glu Glu Arg Gln Arg Gln Gln Arg Arg Glu Glu Arg
 100 105 110
 Arg Gln Gln Asn Leu Arg Ala Arg Ser Arg Glu His Pro Val Val Gly
 115 120 125
 His Pro Asp Pro Ala Leu Pro Pro Ser Gly Val Asn Cys Ser Gly Cys

130	135	140
Gly Ala Glu Leu His Cys Gln Asp Ala Gly Val Pro Gly Tyr Leu Pro		
145	150	155 160
Arg Glu Lys Phe Leu Arg Thr Ala Glu Ala Asp Gly Gly Leu Ala Arg		
	165	170 175
Thr Val Cys Gln Arg Cys Trp Leu Leu Ser His His Arg Arg Ala Leu		
	180	185 190
Arg Leu Gln Val Ser Arg Glu Gln Tyr Leu Glu Leu Val Ser Ala Ala		
	195	200 205
Leu Arg Arg Pro Gly Pro Ser Leu Val Leu Tyr Met Val Asp Leu Leu		
	210	215 220
Asp Leu Pro Asp Ala Leu Leu Pro Asp Leu Pro Ala Leu Val Gly Pro		
	225	230 235 240
Lys Gln Leu Ile Val Leu Gly Asn Lys Val Asp Leu Leu Pro Gln Asp		
	245	250 255
Ala Pro Gly Tyr Arg Gln Arg Leu Arg Glu Arg Leu Trp Glu Asp Cys		
	260	265 270
Ala Arg Ala Gly Leu Leu Leu Ala Pro Gly His Gln Gly Pro Gln Arg		
	275	280 285
Pro Val Lys Asp Glu Pro Gln Asp Gly Glu Asn Pro Asn Pro Pro Asn		
	290	295 300
Trp Ser Arg Thr Val Val Arg Asp Val Arg Leu Ile Ser Ala Lys Thr		
	305	310 315 320
Gly Tyr Gly Val Glu Glu Leu Ile Ser Ala Leu Gln Arg Ser Trp Arg		
	325	330 335
Tyr Arg Gly Asp Val Tyr Leu Val Gly Ala Thr Asn Ala Gly Lys Ser		
	340	345 350
Thr Leu Phe Asn Thr Leu Leu Glu Ser Asp Tyr Cys Thr Ala Lys Gly		
	355	360 365
Ser Asp Ala Ile Asp Arg Ala Thr Ile Ser Pro Trp Pro Gly Thr Thr		
	370	375 380
Leu Asn Leu Leu Lys Phe Pro Ile Cys Asn Pro Thr Pro Tyr Arg Met		
	385	390 395 400
Phe Lys Arg His Gln Arg Leu Lys Lys Asp Ser Thr Gln Ala Glu Glu		
	405	410 415
Asp Leu Ser Glu Gln Glu Gln Asn Gln Leu Asn Val Leu Lys Lys His		
	420	425 430
Gly Tyr Val Val Gly Arg Val Gly Arg Thr Phe Leu Tyr Ser Glu Glu		
	435	440 445

Gln Lys Asp Asn Ile Pro Phe Glu Phe Asp Ala Asp Ser Leu Ala Phe
 450 455 460
 Asp Met Glu Asn Asp Pro Val Met Gly Thr His Lys Ser Thr Lys Gln
 465 470 475 480
 Val Glu Leu Thr Ala Gln Asp Val Lys Asp Ala His Trp Phe Tyr Asp
 485 490 495
 Thr Pro Gly Ile Thr Lys Glu Asn Cys Ile Leu Asn Leu Leu Thr Glu
 500 505 510
 Lys Glu Val Asn Ile Val Leu Pro Thr Gln Ser Ile Val Pro Arg Thr
 515 520 525
 Phe Val Leu Lys Pro Gly Met Val Leu Phe Leu Gly Ala Ile Gly Arg
 530 535 540
 Ile Asp Phe Leu Gln Gly Asn Gln Ser Ala Trp Phe Thr Val Val Ala
 545 550 555 560
 Ser Asn Ile Leu Pro Val His Ile Thr Ser Leu Asp Arg Ala Asp Ala
 565 570 575
 Leu Tyr Gln Lys His Ala Gly His Thr Leu Leu Gln Ile Pro Met Gly
 580 585 590
 Gly Lys Glu Arg Met Ala Gly Phe Pro Pro Leu Val Ala Glu Asp Ile
 595 600 605
 Met Leu Lys Glu Gly Leu Gly Ala Ser Glu Ala Val Ala Asp Ile Lys
 610 615 620
 Phe Ser Ser Ala Gly Trp Val Ser Val Thr Pro Asn Phe Lys Asp Arg
 625 630 635 640
 Leu His Leu Arg Gly Tyr Thr Pro Glu Gly Thr Val Leu Thr Val Arg
 645 650 655
 Pro Pro Leu Leu Pro Tyr Ile Val Asn Ile Lys Gly Gln Arg Ile Lys
 660 665 670
 Lys Ser Val Ala Tyr Lys Thr Lys Lys Pro Pro Ser Leu Met Tyr Asn
 675 680 685
 Val Arg Lys Lys Lys Gly Lys Ile Asn Val
 690 695

<210> 198

<211> 348

<212> PRT

<213> Homo sapiens

<400> 198

Met Asn Met Thr Gln Ala Arg Val Leu Val Ala Ala Val Val Gly Leu
 1 5 10 15

Val Ala Val Leu Leu Tyr Ala Ser Ile His Lys Ile Glu Glu Gly His

20	25	30
Leu Ala Val Tyr Tyr Arg Gly Gly Ala Leu Leu Thr Ser Pro Ser Gly 35 40 45		
Pro Gly Tyr His Ile Met Leu Pro Phe Ile Thr Thr Phe Arg Ser Val 50 55 60		
Gln Thr Thr Leu Gln Thr Asp Glu Val Lys Asn Val Pro Cys Gly Thr 65 70 75 80		
Ser Gly Gly Val Met Ile Tyr Ile Asp Arg Ile Glu Val Val Asn Met 85 90 95		
Leu Ala Pro Tyr Ala Val Phe Asp Ile Val Arg Asn Tyr Thr Ala Asp 100 105 110		
Tyr Asp Lys Thr Leu Ile Phe Asn Lys Ile His His Glu Leu Asn Gln 115 120 125		
Phe Cys Ser Ala His Thr Leu Gln Glu Val Tyr Ile Glu Leu Phe Asp 130 135 140		
Gln Ile Asp Glu Asn Leu Lys Gln Ala Leu Gln Lys Asp Leu Asn Leu 145 150 155 160		
Met Ala Pro Gly Leu Thr Ile Gln Ala Val Arg Val Thr Lys Pro Lys 165 170 175		
Ile Pro Glu Ala Ile Arg Arg Asn Phe Glu Leu Met Glu Ala Glu Lys 180 185 190		
Thr Lys Leu Leu Ile Ala Ala Gln Lys Gln Lys Val Val Glu Lys Glu 195 200 205		
Ala Glu Thr Glu Arg Lys Lys Ala Val Ile Glu Ala Glu Lys Ile Ala 210 215 220		
Gln Val Ala Lys Ile Arg Phe Gln Gln Lys Val Met Glu Lys Glu Thr 225 230 235 240		
Glu Lys Arg Ile Ser Glu Ile Glu Asp Ala Ala Phe Leu Ala Arg Glu 245 250 255		
Lys Ala Lys Ala Asp Ala Glu Tyr Tyr Ala Ala His Lys Tyr Ala Thr 260 265 270		
Ser Asn Lys His Lys Leu Thr Pro Glu Tyr Leu Glu Leu Lys Lys Tyr 275 280 285		
Gln Ala Ile Ala Ser Asn Ser Lys Ile Tyr Phe Gly Ser Asn Ile Pro 290 295 300		
Asn Met Phe Val Asp Ser Ser Cys Ala Leu Lys Tyr Ser Asp Ile Arg 305 310 315 320		
Thr Gly Arg Glu Ser Ser Leu Pro Ser Lys Glu Ala Leu Glu Pro Ser 325 330 335		

Gly Glu Asn Val Ile Gln Asn Lys Glu Ser Thr Gly
 340 345

<210> 199
 <211> 401
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (307)
 <223> Xaa equals any of the naturally occurring L-amino acids

<400> 199
 Met Met Gly Leu Gly Asn Gly Arg Arg Ser Met Lys Ser Pro Pro Leu
 1 5 10 15
 Val Leu Ala Ala Leu Val Ala Cys Ile Ile Val Leu Gly Phe Asn Tyr
 20 25 30
 Trp Ile Ala Ser Ser Arg Ser Val Asp Leu Gln Thr Arg Ile Met Glu
 35 40 45
 Leu Glu Gly Arg Val Arg Arg Arg Ala Ala Glu Arg Gly Ala Val Glu
 50 55 60
 Leu Lys Lys Asn Glu Phe Gln Gly Glu Leu Glu Lys Gln Arg Glu Gln
 65 70 75 80
 Leu Asp Lys Ile Gln Ser Ser His Asn Phe Gln Leu Glu Ser Val Asn
 85 90 95
 Lys Leu Tyr Gln Asp Glu Lys Ala Val Leu Val Asn Asn Ile Thr Thr
 100 105 110
 Gly Glu Arg Leu Ile Arg Val Leu Gln Asp Gln Leu Lys Thr Leu Gln
 115 120 125
 Arg Asn Tyr Gly Arg Leu Gln Gln Asp Val Leu Gln Phe Gln Lys Asn
 130 135 140
 Gln Thr Asn Leu Glu Arg Lys Phe Ser Tyr Asp Leu Ser Gln Cys Ile
 145 150 155 160
 Asn Gln Met Lys Glu Val Lys Glu Gln Cys Glu Glu Arg Ile Glu Glu
 165 170 175
 Val Thr Lys Lys Gly Asn Glu Ala Val Ala Ser Arg Asp Leu Ser Glu
 180 185 190
 Asn Asn Asp Gln Arg Gln Gln Leu Gln Ala Leu Ser Glu Pro Gln Pro
 195 200 205
 Arg Leu Gln Ala Ala Gly Leu Pro His Thr Glu Val Pro Gln Gly Lys
 210 215 220
 Gly Asn Val Leu Gly Asn Ser Lys Ser Gln Thr Pro Ala Pro Ser Ser
 225 230 235 240

Glu Val Val Leu Asp Ser Lys Arg Gln Val Glu Lys Glu Glu Thr Asn
 245 250 255
 Glu Ile Gln Val Val Asn Glu Glu Pro Gln Arg Asp Arg Leu Pro Gln
 260 265 270
 Glu Pro Gly Arg Glu Gln Val Val Glu Asp Arg Pro Val Gly Gly Arg
 275 280 285
 Gly Phe Gly Gly Ala Gly Glu Leu Gly Gln Thr Pro Gln Val Gln Ala
 290 295 300
 Ala Leu Xaa Val Ser Gln Glu Asn Pro Glu Met Glu Gly Pro Glu Arg
 305 310 315 320
 Asp Gln Leu Val Ile Pro Asp Gly Gln Glu Glu Glu Gln Glu Ala Ala
 325 330 335
 Gly Glu Gly Arg Asn Gln Gln Lys Leu Arg Gly Glu Asp Asp Tyr Asn
 340 345 350
 Met Asp Glu Asn Glu Ala Glu Ser Glu Thr Asp Lys Gln Ala Ala Leu
 355 360 365
 Ala Gly Asn Asp Arg Asn Ile Asp Val Phe Asn Val Glu Asp Gln Lys
 370 375 380
 Arg Asp Thr Ile Asn Leu Leu Asp Gln Arg Glu Lys Arg Asn His Thr
 385 390 395 400
 Leu

<210> 200
 <211> 324
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (3)
 <223> Xaa equals any of the naturally occurring L-amino acids

<400> 200
 Met Glu Xaa Ala Lys Val Tyr Val Ala Lys Val Asp Cys Thr Ala His
 1 5 10 15
 Ser Asp Val Cys Ser Ala Gln Gly Val Arg Gly Tyr Pro Thr Leu Lys
 20 25 30
 Leu Phe Lys Pro Gly Gln Glu Ala Val Lys Tyr Gln Gly Pro Arg Asp
 35 40 45
 Phe Gln Thr Leu Glu Asn Trp Met Leu Gln Thr Leu Asn Glu Glu Pro
 50 55 60
 Val Thr Pro Glu Pro Glu Val Glu Pro Pro Ser Ala Pro Glu Leu Lys

65 70 75 80
 Gln Gly Leu Tyr Glu Leu Ser Ala Ser Asn Phe Glu Leu His Val Ala
 85 90 95
 Gln Gly Asp His Phe Ile Lys Phe Phe Ala Pro Trp Cys Gly His Cys
 100 105 110
 Lys Ala Leu Ala Pro Thr Trp Glu Gln Leu Ala Leu Gly Leu Glu His
 115 120 125
 Ser Glu Thr Val Lys Ile Gly Lys Val Asp Cys Thr Gln His Tyr Glu
 130 135 140
 Leu Cys Ser Gly Asn Gln Val Arg Gly Tyr Pro Thr Leu Leu Trp Phe
 145 150 155 160
 Arg Asp Gly Lys Lys Val Asp Gln Tyr Lys Gly Lys Arg Asp Leu Glu
 165 170 175
 Ser Leu Arg Glu Tyr Val Glu Ser Gln Leu Gln Arg Thr Glu Thr Gly
 180 185 190
 Ala Thr Glu Thr Val Thr Pro Ser Glu Ala Pro Val Leu Ala Ala Glu
 195 200 205
 Pro Glu Ala Asp Lys Gly Thr Val Leu Ala Leu Thr Glu Asn Asn Phe
 210 215 220
 Asp Asp Thr Ile Ala Glu Gly Ile Thr Phe Ile Lys Phe Tyr Ala Pro
 225 230 235 240
 Trp Cys Gly His Cys Lys Thr Leu Ala Pro Thr Trp Glu Glu Leu Ser
 245 250 255
 Lys Lys Glu Phe Pro Gly Leu Ala Gly Val Lys Ile Ala Glu Val Asp
 260 265 270
 Cys Thr Ala Glu Arg Asn Ile Cys Ser Lys Tyr Ser Val Arg Gly Tyr
 275 280 285
 Pro Thr Leu Leu Leu Phe Arg Gly Gly Lys Lys Val Ser Glu His Ser
 290 295 300
 Gly Gly Arg Asp Leu Asp Ser Leu His Arg Phe Val Leu Ser Gln Ala
 305 310 315 320
 Lys Asp Glu Leu

<210> 201
 <211> 90
 <212> PRT
 <213> Homo sapiens

<400> 201
 Met Ala Leu Phe Ser Cys Leu Leu Leu Leu Lys Gln Ser Asp Gly Ala
 1 5 10 15

Ser Pro Val Leu Arg Ala Leu Ala Ala Ser Cys Leu Ala Ser Pro Ala
 20 25 30
 Gly Cys Cys Gly Thr Arg Lys Ala Leu Asn Gly Asn Val Gly Glu Lys
 35 40 45
 Val Gly Phe Thr Phe Met Ser Phe Gln Gly Cys Asp Pro Ser Ser Pro
 50 55 60
 Gly Cys Leu Cys Cys Ser Leu Leu Pro Ser Asn Ser Gln Leu Val Phe
 65 70 75 80
 Ile Ser Phe Leu Val Leu Ser Gly Leu Ala
 85 90

<210> 202
 <211> 243
 <212> PRT
 <213> Homo sapiens

<400> 202
 Met Arg Pro Gln Gly Pro Ala Ala Ser Pro Gln Arg Leu Arg Gly Leu
 1 5 10 15
 Leu Leu Leu Leu Leu Gln Leu Pro Ala Pro Ser Ser Ala Ser Glu
 20 25 30
 Ile Pro Lys Gly Lys Gln Lys Ala Gln Leu Arg Gln Arg Glu Val Val
 35 40 45
 Asp Leu Tyr Asn Gly Met Cys Leu Gln Gly Pro Ala Gly Val Pro Gly
 50 55 60
 Arg Asp Gly Ser Pro Gly Ala Asn Gly Ile Pro Gly Thr Pro Gly Ile
 65 70 75 80
 Pro Gly Arg Asp Gly Phe Lys Gly Glu Lys Gly Glu Cys Leu Arg Glu
 85 90 95
 Ser Phe Glu Glu Ser Trp Thr Pro Asn Tyr Lys Gln Cys Ser Trp Ser
 100 105 110
 Ser Leu Asn Tyr Gly Ile Asp Leu Gly Lys Ile Ala Glu Cys Thr Phe
 115 120 125
 Thr Lys Met Arg Ser Asn Ser Ala Leu Arg Val Leu Phe Ser Gly Ser
 130 135 140
 Leu Arg Leu Lys Cys Arg Asn Ala Cys Cys Glu Arg Trp Tyr Phe Thr
 145 150 155 160
 Phe Asn Gly Ala Glu Cys Ser Gly Pro Leu Pro Ile Glu Ala Ile Ile
 165 170 175
 Tyr Leu Asp Gln Gly Ser Pro Glu Met Asn Ser Thr Ile Asn Ile His
 180 185 190

Arg Thr Ser Ser Val Glu Gly Leu Cys Glu Gly Ile Gly Ala Gly Leu
 195 200 205
 Val Asp Val Ala Ile Trp Val Gly Thr Cys Ser Asp Tyr Pro Lys Gly
 210 215 220
 Asp Ala Ser Thr Gly Trp Asn Ser Val Ser Arg Ile Ile Ile Glu Glu
 225 230 235 240
 Leu Pro Lys

<210> 203
 <211> 75
 <212> PRT
 <213> Homo sapiens

<400> 203
 Met Ala Gly Gln Glu Asp Pro Val Gln Arg Glu Ile His Gln Asp Trp
 1 5 10 15
 Ala Asn Arg Glu Tyr Ile Glu Ile Ile Thr Ser Ser Ile Lys Lys Ile
 20 25 30
 Ala Asp Phe Leu Asn Ser Phe Asp Met Ser Cys Arg Ser Arg Leu Ala
 35 40 45
 Thr Leu Asn Glu Lys Leu Thr Ala Leu Glu Arg Arg Ile Glu Tyr Ile
 50 55 60
 Glu Ala Arg Val Thr Lys Gly Glu Thr Leu Thr
 65 70 75

<210> 204
 <211> 248
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (185)
 <223> Xaa equals any of the naturally occurring L-amino acids
 <400> 204
 Met Thr Ser Gln Pro Val Pro Asn Glu Thr Ile Ile Val Leu Pro Ser
 1 5 10 15
 Asn Val Ile Asn Phe Ser Gln Ala Glu Lys Pro Glu Pro Thr Asn Gln
 20 25 30
 Gly Gln Asp Ser Leu Lys Lys His Leu His Ala Glu Ile Lys Val Ile
 35 40 45
 Gly Thr Ile Gln Ile Leu Cys Gly Met Met Val Leu Ser Leu Gly Ile
 50 55 60
 Ile Leu Ala Ser Ala Ser Phe Ser Pro Asn Phe Thr Gln Val Thr Ser

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65              70              75              80
Thr Leu Leu Asn Ser Ala Tyr Pro Phe Ile Gly Pro Phe Phe Phe Ile
      85              90              95
Ile Ser Gly Ser Leu Ser Ile Ala Thr Glu Lys Arg Leu Thr Lys Leu
      100             105             110
Leu Val His Ser Ser Leu Val Gly Ser Ile Leu Ser Ala Leu Ser Ala
      115             120             125
Leu Val Gly Phe Ile Ile Leu Ser Val Lys Gln Ala Thr Leu Asn Pro
      130             135             140
Ala Ser Leu Gln Cys Glu Leu Asp Lys Asn Asn Ile Pro Thr Arg Ser
145             150             155             160
Tyr Val Ser Tyr Phe Tyr His Asp Ser Leu Tyr Thr Thr Asp Cys Tyr
      165             170             175
Thr Ala Lys Ala Ser Leu Ala Gly Xaa Leu Ser Leu Met Leu Ile Cys
      180             185             190
Thr Leu Leu Glu Phe Cys Leu Ala Val Leu Thr Ala Val Leu Arg Trp
      195             200             205
Lys Gln Ala Tyr Ser Asp Phe Pro Gly Ser Val Leu Phe Leu Pro His
      210             215             220
Ser Tyr Ile Gly Asn Ser Gly Met Ser Ser Lys Met Thr His Asp Cys
225             230             235             240
Gly Tyr Glu Glu Leu Leu Thr Ser
      245

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<210> 205

<211> 168

<212> PKT

<213> Homo sapiens

<220>

<221> SITE

<222> (83)

<223> Xaa equals any of the naturally occurring L-amino acids

<400> 205

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Met Pro Leu Leu Arg Gly Leu Leu Trp Leu Gln Val Leu Cys Ala Gly
  1              5              10              15
Pro Leu His Thr Glu Ala Val Val Leu Leu Val Pro Ser Asp Asp Gly
      20              25              30
Arg Ala Phe Leu Leu Arg Ser Arg Leu Leu His Pro Glu Ala His Val
      35              40              45
Pro Pro Ala Ala Asp Arg Gly Ala Ser Leu Gln Cys Val Leu His Gln
      50              55              60

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Ala Ala Pro Lys Ser Arg Pro Arg Ser Pro Ala Ala Gly Ala Ala Leu
 65 70 75 80
 Leu His Xaa Pro Arg Arg Thr Gly Asp Glu Pro Cys Arg Glu Phe His
 85 90 95
 Gly Asn Gly Phe Pro Gly Pro Thr Gln Leu Thr Pro Gly Glu Cys Gly
 100 105 110
 Leu Pro Ala Pro Ser Ser Leu Leu Gln His Ala Ser Ala Pro Val Arg
 115 120 125
 Thr Gly Ser Glu Gly Gln Val Val Gly Cys Pro Arg Ala Arg Gly Glu
 130 135 140
 Thr Gly Glu Gly Leu Ser Leu Ala Phe Leu Ser Ser Leu Met Phe Thr
 145 150 155 160
 Ser Arg Asn Gly Leu Val Gly Cys
 165

<210> 206
 <211> 218
 <212> PRT
 <213> Homo sapiens

<400> 206
 Met Gly Ser Ala Ala Leu Glu Ile Leu Gly Leu Val Leu Cys Leu Val
 1 5 10 15
 Gly Trp Gly Gly Leu Ile Leu Ala Cys Gly Leu Pro Met Trp Gln Val
 20 25 30
 Thr Ala Phe Leu Asp His Asn Ile Val Thr Ala Gln Thr Thr Trp Lys
 35 40 45
 Gly Leu Trp Met Ser Cys Val Val Gln Ser Thr Gly His Met Gln Cys
 50 55 60
 Lys Val Tyr Asp Ser Val Leu Ala Leu Ser Thr Glu Val Gln Ala Ala
 65 70 75 80
 Arg Ala Leu Thr Val Ser Ala Val Leu Leu Ala Phe Val Ala Leu Phe
 85 90 95
 Val Thr Leu Ala Gly Ala Gln Cys Thr Thr Cys Val Ala Pro Gly Pro
 100 105 110
 Ala Lys Ala Arg Val Ala Leu Thr Gly Gly Val Leu Tyr Leu Phe Cys
 115 120 125
 Gly Leu Leu Ala Leu Val Pro Leu Cys Trp Phe Ala Asn Ile Val Val
 130 135 140
 Arg Glu Phe Tyr Asp Pro Ser Val Pro Val Ser Gln Lys Tyr Glu Leu
 145 150 155 160
 Gly Ala Ala Leu Tyr Ile Gly Trp Ala Ala Thr Ala Leu Leu Met Val

165 170 175
 Gly Gly Cys Leu Leu Cys Cys Gly Ala Trp Val Cys Thr Gly Arg Pro
 180 185 190
 Asp Leu Ser Phe Pro Val Lys Tyr Ser Ala Pro Arg Arg Pro Thr Ala
 195 200 205
 Thr Gly Asp Tyr Asp Lys Lys Asn Tyr Val
 210 215

<210> 207
 <211> 73
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (73)
 <223> Xaa equals stop translation

<400> 207
 Met Thr Ser Tyr Ile Leu Ile Ser Phe Val Leu Leu Ile Gly Val Gly
 1 5 10 15
 Cys Ile Glu Lys Asp Gln Ser Cys Pro Val Phe Gly Gly Arg Lys Arg
 20 25 30
 Leu His Leu Leu Phe Val Gly Gly Gln Leu Arg Gln Val Arg Met Leu
 35 40 45
 Arg Gly Glu Leu Ser Cys Ala Cys Tyr Arg Pro His Val Gln Ala Leu
 50 55 60
 Gln Leu Gly Gly Cys Thr Cys Phe Xaa
 65 70

<210> 208
 <211> 348
 <212> PRT
 <213> Homo sapiens

<400> 208
 Met Leu Cys Pro Trp Arg Thr Ala Asn Leu Gly Leu Leu Ile Leu
 1 5 10 15
 Thr Ile Phe Leu Val Ala Glu Ala Glu Gly Ala Ala Gln Pro Asn Asn
 20 25 30
 Ser Leu Met Leu Gln Thr Ser Lys Glu Asn His Ala Leu Ala Ser Ser
 35 40 45
 Ser Leu Cys Met Asp Glu Lys Gln Ile Thr Gln Asn Tyr Ser Lys Val
 50 55 60
 Leu Ala Glu Val Asn Thr Ser Trp Pro Val Lys Met Ala Thr Asn Ala
 65 70 75 80

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<210> 209
<211> 73
<212> PRT
<213> Homo sapiens
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<220>

<221> SITE
 <222> (73)
 <223> Xaa equals stop translation

 <400> 209
 Met Ala Arg Gly Cys Val Cys Ser Leu Cys Ala Ser Val Cys Ile Phe
 1 5 10 15
 Leu Ser Ser Leu Phe Pro Leu Leu Pro Ser Val His Ser Val Asn Ile
 20 25 30
 Ile Ser Cys Leu Leu Leu Ser Lys Cys Phe Glu Gly Leu Glu Leu Met
 35 40 45
 Cys Glu His Leu Tyr Gln Leu Ser Gln Leu His Val Leu His His Ile
 50 55 60
 Phe Ser Tyr Leu Leu Cys Thr Pro Xaa
 65 70

 <210> 210
 <211> 608
 <212> PRT
 <213> Homo sapiens

 <220>
 <221> SITE
 <222> (265)
 <223> Xaa equals any of the naturally occurring L-amino acids

 <220>
 <221> SITE
 <222> (597)
 <223> Xaa equals any of the naturally occurring L-amino acids

 <400> 210
 Met Val Gly Thr Lys Leu Arg Gln Thr Lys Asp Ala Leu Phe Thr Ile
 1 5 10 15
 Leu His Asp Leu Arg Pro Gln Asp Arg Phe Ser Ile Ile Gly Phe Ser
 20 25 30
 Asn Arg Ile Lys Val Trp Lys Asp His Leu Ile Ser Val Thr Pro Asp
 35 40 45
 Ser Ile Arg Asp Gly Lys Val Tyr Ile His His Met Ser Pro Thr Gly
 50 55 60
 Gly Thr Asp Ile Asn Gly Val Leu Gln Arg Ala Ile Arg Leu Leu Asn
 65 70 75 80
 Lys Tyr Val Ala His Ser Gly Ile Gly Asp Arg Ser Val Ser Leu Ile
 85 90 95
 Val Phe Leu Thr Asp Gly Lys Pro Thr Val Gly Glu Thr His Thr Leu
 100 105 110
 Lys Ile Leu Asn Asn Thr Arg Glu Ala Ala Arg Gly Gln Val Cys Ile

115	120	125
Phe Thr Ile Gly Ile Gly Asn Asp Val Asp Phe Arg Leu Leu Glu Lys 130 135 140		
Leu Ser Leu Glu Asn Cys Gly Leu Thr Arg Arg Val His Glu Glu Glu 145 150 155 160		
Asp Ala Gly Ser Gln Leu Ile Gly Phe Tyr Asp Glu Ile Arg Thr Pro 165 170 175		
Leu Leu Ser Asp Ile Arg Ile Asp Tyr Pro Pro Ser Ser Val Val Gln 180 185 190		
Ala Thr Lys Thr Leu Phe Pro Asn Tyr Phe Asn Gly Ser Glu Ile Ile 195 200 205		
Ile Ala Gly Lys Leu Val Asp Arg Lys Leu Asp His Leu His Val Glu 210 215 220		
Val Thr Ala Ser Asn Ser Lys Lys Phe Ile Ile Leu Lys Thr Asp Val 225 230 235 240		
Pro Val Arg Pro Gln Lys Ala Gly Lys Asp Val Thr Gly Ser Pro Arg 245 250 255		
Pro Gly Gly Asp Gly Glu Gly Asp Xaa Asn His Ile Glu Arg Leu Trp 260 265 270		
Ser Tyr Leu Thr Thr Lys Glu Leu Leu Ser Ser Trp Leu Gln Ser Asp 275 280 285		
Asp Glu Pro Glu Lys Glu Arg Leu Arg Gln Arg Ala Gln Ala Leu Ala 290 295 300		
Val Ser Tyr Arg Phe Leu Thr Pro Phe Thr Ser Met Lys Leu Arg Gly 305 310 315 320		
Pro Val Pro Arg Met Asp Gly Leu Glu Glu Ala His Gly Met Ser Ala 325 330 335		
Ala Met Gly Pro Glu Pro Val Val Gln Ser Val Arg Gly Ala Gly Thr 340 345 350		
Gln Pro Gly Pro Leu Leu Lys Lys Pro Tyr Gln Pro Arg Ile Lys Ile 355 360 365		
Ser Lys Thr Ser Val Asp Gly Asp Pro His Phe Val Val Asp Phe Pro 370 375 380		
Leu Ser Arg Leu Thr Val Cys Phe Asn Ile Asp Gly Gln Pro Gly Asp 385 390 395 400		
Ile Leu Arg Leu Val Ser Asp His Arg Asp Ser Gly Val Thr Val Asn 405 410 415		
Gly Glu Leu Ile Gly Ala Pro Ala Pro Pro Asn Gly His Lys Lys Gln 420 425 430		

Arg Thr Tyr Leu Arg Thr Ile Thr Ile Leu Ile Asn Lys Pro Glu Arg
 435 440 445
 Ser Tyr Leu Glu Ile Thr Pro Ser Arg Val Ile Leu Asp Gly Gly Asp
 450 455 460
 Arg Leu Val Leu Pro Cys Asn Gln Ser Val Val Val Gly Ser Trp Gly
 465 470 475 480
 Leu Glu Val Ser Val Ser Ala Asn Ala Asn Val Thr Val Thr Ile Gln
 485 490 495
 Gly Ser Ile Ala Phe Val Ile Leu Ile His Leu Tyr Lys Lys Pro Ala
 500 505 510
 Pro Phe Gln Arg His His Leu Gly Phe Tyr Ile Ala Asn Ser Glu Gly
 515 520 525
 Leu Ser Ser Asn Cys His Gly Leu Leu Gly Gln Phe Leu Asn Gln Asp
 530 535 540
 Ala Arg Leu Thr Glu Asp Pro Ala Gly Pro Ser Gln Asn Leu Thr His
 545 550 555 560
 Pro Leu Leu Leu Gln Val Gly Glu Gly Pro Glu Ala Val Leu Thr Val
 565 570 575
 Lys Gly His Gln Val Pro Val Val Trp Lys Gln Arg Lys Ile Tyr Asn
 580 585 590
 Gly Glu Glu Gln Xaa Asp Cys Trp Phe Ala Arg Asn Met Pro Pro Asn
 595 600 605

<210> 211
 <211> 252
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (252)
 <223> Xaa equals stop translation

<400> 211
 Met Ala Pro Ala Ser Arg Leu Leu Ala Leu Trp Ala Leu Ala Ala Val
 1 5 10 15
 Ala Leu Pro Gly Ser Gly Ala Glu Gly Asp Gly Gly Trp Arg Pro Gly
 20 25 30
 Gly Pro Gly Ala Val Ala Glu Glu Glu Arg Cys Thr Val Glu Arg Arg
 35 40 45
 Ala Asp Leu Thr Tyr Ala Glu Phe Val Gln Gln Tyr Ala Phe Val Arg
 50 55 60

Pro Val Ile Leu Gln Gly Leu Thr Asp Asn Ser Arg Phe Arg Ala Leu
 65 70 75 80
 Cys Ser Arg Asp Arg Leu Leu Ala Ser Phe Gly Asp Arg Val Val Arg
 85 90 95
 Leu Ser Thr Ala Asn Thr Tyr Ser Tyr His Lys Val Asp Leu Pro Phe
 100 105 110
 Gln Glu Tyr Val Glu Gln Leu Leu His Pro Gln Asp Pro Thr Ser Leu
 115 120 125
 Gly Asn Asp Thr Leu Tyr Phe Phe Gly Asp Asn Asn Phe Thr Glu Trp
 130 135 140
 Ala Ser Leu Phe Arg His Tyr Ser Pro Pro Pro Phe Gly Leu Leu Gly
 145 150 155 160
 Thr Ala Pro Ala Tyr Ser Phe Gly Ile Ala Gly Ala Gly Ser Gly Val
 165 170 175
 Pro Phe His Trp His Gly Pro Gly Tyr Ser Glu Val Ile Tyr Gly Arg
 180 185 190
 Lys Arg Trp Phe Leu Tyr Pro Pro Glu Lys Thr Pro Glu Phe His Pro
 195 200 205
 Asn Lys Thr Thr Leu Ala Trp Leu Arg Asp Thr Tyr Pro Ala Cys Thr
 210 215 220
 Val Cys Thr Ala Leu Glu Cys Thr Ile Arg Ala Gly Glu Val Leu Thr
 225 230 235 240
 Ser Arg Pro Leu Val Ala Cys Tyr Ala Gln Pro Xaa
 245 250

<210> 212
 <211> 226
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (226)
 <223> Xaa equals stop translation

<400> 212
 Met Lys Glu Ile Pro Ala Leu Leu His Leu Pro Val Leu Ile Ile Met
 1 5 10 15
 Ala Leu Ala Ile Leu Ser Phe Cys Tyr Gly Ala Gly Lys Ser Val His
 20 25 30
 Val Leu Arg His Ile Gly Gly Pro Glu Arg Glu Pro Pro Gln Ala Leu
 35 40 45
 Arg Pro Arg Asp Arg Arg Arg Gln Glu Glu Ile Asp Tyr Arg Pro Asp

<210> 214
 <211> 172
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (172)
 <223> Xaa equals stop translation

<400> 214
 Met Trp Leu Trp Ala Val Ser Pro Val Arg Pro Arg Thr Cys Leu Pro
 1 5 10 15
 Pro Cys Pro Arg Leu Trp Leu Trp Ile Ser Met Thr Leu Val Pro Ser
 20 25 30
 Ser Ser Ala Trp Lys Ser His Gly Ala Pro Ser Thr Arg Met Thr Ser
 35 40 45
 Pro Gln Leu Leu Leu Ser Thr Arg Pro Pro Gln Ser Pro Ser Ala
 50 55 60
 Ser Pro Pro Ile Ala Arg Ala His Arg Thr His Pro His Phe Gly Asn
 65 70 75 80
 Arg Leu Ser Ile Thr Cys Cys Asp Gly Arg Arg Ser Trp Arg Met Gly
 85 90 95
 Gln His Gly Pro Cys His Leu Asn Leu Gln Thr Thr His Pro Ala His
 100 105 110
 Ser Ser Gln Ala Leu Pro Ala Thr His Gln Pro Leu Gly Pro Trp Cys
 115 120 125
 Ser Ser Pro Ser Pro Phe Pro Ser Lys Leu Pro Ser Ala Gly Leu Arg
 130 135 140
 Pro Pro Ala Leu Gly Pro Trp Met Arg Arg Gly Pro Trp Pro Gln Ser
 145 150 155 160
 Trp Gln Met Gly Met His Pro Thr Val Gly Leu Xaa
 165 170

<210> 215
 <211> 48
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (48)
 <223> Xaa equals stop translation

<400> 215
 Met Trp Leu Leu Ile Ile Phe Cys Lys Ser Ala Ser Ala Ser Val Leu
 1 5 10 15

Cys Trp Ile Lys Lys Phe His Pro Val Phe Gln Glu Ser Leu Leu Tyr
 20 25 30
 Leu Val Gln Glu Gly Ser Leu Cys Tyr Val Gln Gln Lys Val Pro Xaa
 35 40 45

<210> 216
 <211> 139
 <212> PRT
 <213> Homo sapiens

<400> 216
 Met Glu Ala Val Val Phe Val Phe Ser Leu Leu Asp Cys Cys Ala Leu
 1 5 10 15
 Ile Phe Leu Ser Val Tyr Phe Ile Ile Thr Leu Ser Asp Leu Glu Cys
 20 25 30
 Asp Tyr Ile Asn Ala Arg Ser Cys Cys Ser Lys Leu Asn Lys Trp Val
 35 40 45
 Ile Pro Glu Leu Ile Gly His Thr Ile Val Thr Val Leu Leu Leu Met
 50 55 60
 Ser Leu His Trp Phe Ile Phe Leu Leu Asn Leu Pro Val Ala Thr Trp
 65 70 75 80
 Asn Ile Tyr Arg Tyr Ile Met Val Pro Ser Gly Asn Met Gly Val Phe
 85 90 95
 Asp Pro Thr Glu Ile His Asn Arg Gly Gln Leu Lys Ser His Met Lys
 100 105 110
 Glu Ala Met Ile Lys Leu Gly Phe His Leu Leu Cys Phe Phe Met Tyr
 115 120 125
 Leu Tyr Ser Met Ile Leu Ala Leu Ile Asn Asp
 130 135

<210> 217
 <211> 41
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (41)
 <223> Xaa equals stop translation

<400> 217
 Met Ser Gly Ser Ser Leu Pro Ser Ala Leu Ala Leu Ser Leu Leu Leu
 1 5 10 15

Val Ser Gly Ser Leu Leu Pro Gly Pro Gly Ala Ala Gln Asn Val Arg
20 25 30

Val Gln Ser Gly Gln Asp Gln Lys Xaa
35 40

<210> 218
<211> 52
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (52)
<223> Xaa equals stop translation

<400> 218
Met Pro Ser His Ile Arg Ala His Leu Phe Leu Leu Leu Phe Phe Leu
1 5 10 15
Phe Ile Tyr Gln Gly Ile Ser Ser Ile Ser Gln Ala Ser Gly Leu Thr
20 25 30
Leu Lys Thr Gln Asn Glu Lys Asp Ile Gln Val Ser Ile Leu Lys Glu
35 40 45
Phe Val Val Xaa
50

<210> 219
<211> 49
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (49)
<223> Xaa equals stop translation

<400> 219
Met Cys Ile Tyr Gln Ser Glu Gln Met Leu Ala Leu Leu Val Leu
1 5 10 15
Val Phe Cys Ile Ser Leu Leu Val Leu Val Cys Trp Gly Ser His Asn
20 25 30
Lys Val Pro Gln Lys Phe Ile Phe Ser Gln Phe Trp Gly Leu Glu Asp
35 40 45
Xaa

<210> 220
<211> 42
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (42)
<223> Xaa equals stop translation

<400> 220
Met Ala Val Pro Leu Phe Leu Tyr Ile Phe Thr Leu Leu Pro Leu Leu
1 5 10 15
Pro Phe Leu Leu Ser Leu Cys Phe Ser Pro Leu Thr Val Lys Arg Ser
20 25 30
Ser Ser Ser Glu Ser Lys Ser Ser Leu Xaa
35 40

<210> 221
<211> 41
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (41)
<223> Xaa equals stop translation

<400> 221
Met Gly Met Leu Leu Ala Phe Trp Leu Pro Gly Ala Ser Trp Gln Glu
1 5 10 15
Ala Gly Pro Arg Ala Ser Thr Gln Arg Met Arg Thr Gln Thr Gln Met
20 25 30
Ser Thr Arg Lys Pro Lys Pro Ala Xaa
35 40

<210> 222
<211> 43
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (43)
<223> Xaa equals stop translation

<400> 222
Met Glu Pro Ala Met Val Leu Lys Phe Leu Ser Ser Leu Pro Glu Asn
1 5 10 15
Leu Phe Leu Pro Ser Leu Leu Phe Phe Ala Trp Leu Cys Trp Asn Met
20 25 30
Val Cys Gly Ser Pro Val Ser Cys Pro Tyr Xaa
35 40

<210> 223
<211> 204
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (204)
<223> Xaa equals stop translation

<400> 223
Met Gln Leu Gly Ser Val Leu Leu Thr Arg Cys Pro Phe Trp Gly Cys
1 5 10 15
Phe Ser Gln Leu Met Leu Tyr Ala Glu Arg Ala Glu Ala Arg Arg Lys
20 25 30
Pro Asp Ile Pro Val Pro Tyr Leu Tyr Phe Asp Met Gly Ala Ala Val
35 40 45
Leu Cys Ala Ser Phe Met Ser Phe Gly Val Lys Arg Arg Trp Phe Ala
50 55 60
Leu Gly Ala Ala Leu Gln Leu Ala Ile Ser Thr Tyr Ala Ala Tyr Ile
65 70 75 80
Gly Gly Tyr Val His Tyr Gly Asp Trp Leu Lys Val Arg Met Tyr Ser
85 90 95
Arg Thr Val Ala Ile Ile Gly Gly Phe Leu Val Leu Ala Ser Gly Ala
100 105 110
Gly Glu Leu Tyr Arg Arg Lys Pro Arg Ser Arg Ser Leu Gln Ser Thr
115 120 125
Gly Gln Val Phe Leu Gly Ile Tyr Leu Ile Cys Val Ala Tyr Ser Leu
130 135 140
Gln His Ser Lys Glu Asp Arg Leu Ala Tyr Leu Asn His Leu Pro Gly
145 150 155 160
Gly Glu Leu Met Ile Gln Leu Phe Phe Val Leu Tyr Gly Ile Leu Ala
165 170 175
Pro Gly Leu Ser Val Arg Leu Leu Arg Asp Pro Arg Cys Pro Asp Pro
180 185 190
Gly Cys Thr Ala Ala Pro Cys His Ala Ala His Xaa
195 200

<210> 224
<211> 43
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (43)

<223> Xaa equals stop translation

<400> 224

Met Arg Val Arg Ile Gly Leu Thr Leu Leu Leu Cys Ala Val Leu Leu
1 5 10 15

Ser Leu Ala Ser Ala Ser Ser Asp Glu Glu Gly Ser Gln Asp Glu Ser
20 25 30

Leu Gly Phe Gln Asp Tyr Phe Asp Ile Arg Xaa
35 40

<210> 225

<211> 156

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (156)

<223> Xaa equals stop translation

<400> 225

Met Ala Arg Gly Ser Leu Arg Arg Leu Leu Arg Leu Leu Val Leu Gly
1 5 10 15

Leu Trp Leu Ala Leu Leu Arg Ser Val Ala Gly Glu Gln Ala Pro Gly
20 25 30

Thr Ala Pro Cys Ser Arg Gly Ser Ser Trp Ser Ala Asp Leu Asp Lys
35 40 45

Cys Met Asp Cys Ser Thr Ser Cys Pro Leu Pro Ala Ala Leu Ala His
50 55 60

Pro Trp Gly Arg Ser Glu Pro Asp Leu Arg Ala Gly Ala Ala Phe Trp
65 70 75 80

Leu Phe Gly Leu Glu Thr Met Pro Gln Glu Arg Glu Val His His Pro
85 90 95

His Arg Gly Asp Arg Arg Arg Gly Leu Pro Ser Cys Gly Ala Asp Pro
100 105 110

Val Thr Met Cys Pro Leu Pro Ala Gly Ala Arg Pro Leu Ile Ile His
115 120 125

Ser Ser Ile Leu Glu Pro Val Ser Ala Ser Gln Thr Arg Arg Glu Pro
130 135 140

Ser Ser Ser Asn His Lys Gly Gly Gly Gly Arg Xaa
145 150 155

<210> 226

<211> 74

<212> PRT

<213> Homo sapiens

<220>
 <221> SITE
 <222> (38)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (48)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (54)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (55)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (68)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (74)
 <223> Xaa equals stop translation

<400> 226
 Met Phe Tyr Lys Leu Thr Leu Ile Leu Cys Glu Leu Ser Val Ala Gly
 1 5 10 15
 Val Thr Gln Ala Ala Ser Gln Arg Pro Leu Gln Arg Leu Pro Arg His
 20 25 30
 Ile Cys Ser Gln Arg Xaa Pro Pro Gly Arg Cys Leu Leu Lys Ala Xaa
 35 40 45
 Leu Gln Thr Thr Trp Xaa Xaa Pro Asp Lys Pro Ile Pro Arg Leu Ser
 50 55 60
 Pro Pro Leu Xaa Ser Asp Pro Lys Arg Xaa
 65 70

<210> 227
 <211> 167
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (167)
 <223> Xaa equals stop translation

<400> 227

Met Gly Ser Arg Phe Leu Leu Val Leu Leu Ser Gly Leu Thr Val Leu
 1 5 10 15
 Leu Ala Leu Pro Gly Ser Glu Ala Lys Asn Ser Gly Ala Ser Cys Pro
 20 25 30
 Pro Cys Pro Lys Tyr Ala Ser Cys His Asn Ser Thr His Cys Thr Cys
 35 40 45
 Glu Asp Gly Phe Arg Ala Arg Ser Gly Arg Thr Tyr Phe His Asp Ser
 50 55 60
 Ser Glu Lys Cys Glu Asp Ile Asn Glu Cys Glu Thr Gly Leu Ala Lys
 65 70 75 80
 Cys Lys Tyr Lys Ala Tyr Cys Arg Asn Lys Val Gly Gly Tyr Ile Cys
 85 90 95
 Ser Cys Leu Val Lys Tyr Thr Leu Phe Asn Phe Leu Ala Gly Ile Ile
 100 105 110
 Asp Tyr Asp His Pro Asp Cys Tyr Glu Asn Asn Ser Gln Gly Thr Thr
 115 120 125
 Gln Ser Asn Val Asp Ile Trp Val Ser Gly Val Lys Pro Gly Phe Gly
 130 135 140
 Lys Gln Leu Val Arg Ile Thr Met Pro Phe Ser Tyr Pro Asn Ile Asn
 145 150 155 160
 Met Ser Ser Cys Asp Phe Xaa
 165

<210> 228

<211> 71

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (71)

<223> Xaa equals stop translation

<400> 228

Met Lys Pro Lys His Leu Glu Trp Cys Leu Ala His Ser Trp Cys Val
 1 5 10 15
 Ile Trp Leu Ser Phe Val Ser Pro Pro Thr Ser His Leu Glu Cys Asp
 20 25 30
 Gly Phe Pro Gly Ser Leu Leu Pro Pro Cys Glu Glu Gly Arg Cys Phe
 35 40 45
 Pro Phe Thr Phe His His His Asp Cys His Gly Cys Ser Pro Leu Gln
 50 55 60
 Ser Ser Pro Gly Gln His Xaa
 65 70

<210> 229
 <211> 273
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (273)
 <223> Xaa equals stop translation

<400> 229
 Met Cys Cys Trp Pro Leu Leu Leu Leu Trp Gly Leu Leu Pro Gly Thr
 1 5 10 15
 Ala Ala Gly Gly Ser Gly Arg Thr Tyr Pro His Arg Thr Leu Leu Asp
 20 25 30
 Ser Glu Gly Lys Tyr Trp Leu Gly Trp Ser Gln Arg Gly Ser Gln Ile
 35 40 45
 Ala Phe Arg Leu Gln Val Arg Thr Ala Gly Tyr Val Gly Phe Gly Phe
 50 55 60
 Ser Pro Thr Gly Ala Met Ala Ser Ala Asp Ile Val Val Gly Gly Val
 65 70 75 80
 Ala His Gly Arg Pro Tyr Leu Gln Asp Tyr Phe Thr Asn Ala Asn Arg
 85 90 95
 Glu Leu Lys Lys Asp Ala Gln Gln Asp Tyr His Leu Glu Tyr Ala Met
 100 105 110
 Glu Asn Ser Thr His Thr Ile Ile Glu Phe Thr Arg Glu Leu His Thr
 115 120 125
 Cys Asp Ile Asn Asp Lys Ser Ile Thr Asp Ser Thr Val Arg Val Ile
 130 135 140
 Trp Ala Tyr His His Glu Asp Ala Gly Glu Ala Gly Pro Lys Tyr His
 145 150 155 160
 Asp Ser Asn Arg Gly Thr Lys Ser Leu Arg Leu Leu Asn Pro Glu Lys
 165 170 175
 Thr Ser Val Leu Ser Thr Ala Leu Pro Tyr Phe Asp Leu Val Asn Gln
 180 185 190
 Asp Val Pro Ile Pro Asn Lys Asp Thr Thr Tyr Trp Cys Gln Met Phe
 195 200 205
 Lys Ile Pro Val Phe Gln Glu Lys His His Val Ile Lys Val Glu Pro
 210 215 220
 Val Ile Gln Arg Gly His Glu Ser Leu Val His His Ile Leu Leu Tyr
 225 230 235 240
 Gln Cys Ser Asn Asn Phe Asn Asp Ser Val Pro Gly Ile Arg Ala Arg

245 250 255
 Ile Ala Ile Thr Pro Thr Cys Pro Met His Ser Ser Pro Val Lys Leu
 260 265 270

Xaa

<210> 230
 <211> 82
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (82)
 <223> Xaa equals stop translation

<400> 230
 Met Arg Pro Gly Thr Ala Leu Gln Ala Val Leu Leu Ala Val Leu Leu
 1 5 10 15
 Val Gly Leu Arg Ala Ala Thr Gly Arg Leu Leu Ser Gly Gln Pro Val
 20 25 30
 Cys Arg Gly Gly Thr Gln Arg Pro Cys Tyr Lys Val Ile Tyr Phe His
 35 40 45
 Asp Thr Ser Arg Arg Leu Asn Phe Glu Glu Ala Lys Glu Ala Cys Arg
 50 55 60
 Arg Gly Trp Arg Pro Ala Ser Gln His Arg Val Leu Lys Met Asn Arg
 65 70 75 80

Asn Xaa

<210> 231
 <211> 71
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (38)
 <223> Xaa equals any of the naturally occurring L-amino acids

<400> 231
 Met Ser Pro Leu Ser Ala Ala Arg Ala Ala Leu Arg Val Tyr Ala Val
 1 5 10 15
 Gly Ala Ala Val Ile Leu Ala Gln Leu Leu Arg Arg Cys Arg Gly Gly
 20 25 30
 Phe Leu Glu Pro Val Xaa Pro Pro Arg Pro Asp Arg Val Ala Ile Val
 35 40 45

Thr Gly Gly Thr Asp Gly Ile Gly Tyr Ser Thr Ala Asn Ile Trp Arg
50 55 60

Asp Leu Ala Cys Met Leu Ser
65 70

<210> 232

<211> 225

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (5)

<223> Xaa equals any of the naturally occurring L-amino acids

<400> 232

His Glu Arg Ala Xaa Gly Pro Ser Arg Gly His Gly Glu Leu Leu Ser
1 5 10 15

Cys Val Leu Gly Pro Arg Leu Tyr Lys Ile Tyr Arg Glu Arg Asp Ser
20 25 30

Glu Arg Ala Pro Ala Ser Val Pro Glu Thr Pro Thr Ala Val Thr Ala
35 40 45

Pro His Ser Ser Ser Trp Asp Thr Tyr Tyr Gln Pro Arg Ala Leu Glu
50 55 60

Lys His Ala Asp Ser Ile Leu Ala Leu Ala Ser Val Phe Trp Ser Ile
65 70 75 80

Ser Tyr Tyr Ser Ser Pro Phe Ala Phe Phe Tyr Leu Tyr Arg Lys Gly
85 90 95

Tyr Leu Ser Leu Ser Lys Val Val Pro Phe Ser His Tyr Ala Gly Thr
100 105 110

Leu Leu Leu Leu Ala Gly Val Ala Cys Ser Glu Ala Leu Ala Ala
115 120 125

Gly Pro Thr Pro Ser Thr Gly Ser Ser Ser Pro Ser Trp Lys Gln His
130 135 140

Ile Gly Thr Ser Leu Gln Lys Thr Arg Gly Ser Leu Pro Thr Thr Thr
145 150 155 160

Leu Thr Ser Gly Ala Gly Gln Ser Thr Ser Thr Gly Lys Asn Pro Ala
165 170 175

Ala Gly Arg Ser Leu Glu Gly Ala Leu Pro Ala Gly Val Trp Pro Cys
180 185 190

Phe Ala Gln Ser Pro Cys Thr Gly Gly Gln Gln Thr Pro Ser Ser Thr
195 200 205

Gly Leu Arg Ser Cys Leu Val Arg Ser Pro Ala Thr Trp Trp Arg Thr
210 215 220

Pro
225

<210> 233
<211> 314
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (147)
<223> Xaa equals any of the naturally occurring L-amino acids

<220>
<221> SITE
<222> (211)
<223> Xaa equals any of the naturally occurring L-amino acids

<400> 233

Met	Leu	Pro	Ala	Arg	Leu	Pro	Phe	Arg	Leu	Leu	Ser	Leu	Phe	Leu	Arg	1	5	10	15
Gly	Ser	Ala	Pro	Thr	Ala	Ala	Arg	His	Gly	Leu	Arg	Glu	Pro	Leu	Leu	20	25	30	
Glu	Arg	Arg	Cys	Ala	Ala	Ala	Ser	Ser	Phe	Gln	His	Ser	Ser	Ser	Leu	35	40	45	
Gly	Arg	Glu	Leu	Pro	Tyr	Asp	Pro	Val	Asp	Thr	Glu	Gly	Phe	Gly	Glu	50	55	60	
Gly	Gly	Asp	Met	Gln	Glu	Arg	Phe	Leu	Phe	Pro	Glu	Tyr	Ile	Leu	Asp	65	70	75	80
Pro	Glu	Pro	Gln	Pro	Thr	Arg	Glu	Lys	Gln	Leu	Gln	Glu	Leu	Gln	Gln	85	90	95	
Gln	Gln	Glu	Glu	Glu	Glu	Arg	Gln	Arg	Gln	Gln	Arg	Arg	Glu	Glu	Arg	100	105	110	
Arg	Gln	Gln	Asn	Leu	Arg	Ala	Arg	Ser	Arg	Glu	His	Pro	Val	Val	Gly	115	120	125	
His	Pro	Asp	Pro	Ala	Leu	Pro	Pro	Ser	Gly	Val	Asn	Cys	Ser	Gly	Cys	130	135	140	
Gly	Ala	Xaa	Leu	His	Cys	Gln	Asp	Ala	Gly	Val	Pro	Gly	Tyr	Leu	Pro	145	150	155	160
Arg	Glu	Lys	Phe	Leu	Arg	Thr	Ala	Glu	Ala	Asp	Gly	Gly	Leu	Ala	Arg	165	170	175	
Thr	Val	Cys	Gln	Arg	Cys	Trp	Leu	Leu	Ser	His	His	Arg	Arg	Ala	Leu	180	185	190	
Arg	Leu	Gln	Val	Ser	Arg	Glu	Gln	Tyr	Leu	Glu	Leu	Val	Ser	Ala	Ala	195	200	205	

Leu Arg Xaa Pro Gly Pro Ser Leu Val Leu Tyr Met Val Asp Leu Leu
 210 215 220
 Asp Leu Pro Asp Ala Leu Leu Pro Asp Leu Pro Ala Leu Val Gly Pro
 225 230 235 240
 Lys Gln Leu Ile Val Leu Gly Asn Lys Val Asp Leu Leu Pro Gln Asp
 245 250 255
 Ala Pro Gly Tyr Arg Gln Arg Leu Arg Glu Arg Leu Trp Glu Asp Cys
 260 265 270
 Ala Arg Ala Gly Leu Leu Leu Ala Pro Gly Thr Lys Gly His Ser Ala
 275 280 285
 Pro Ser Arg Thr Ser His Arg Thr Gly Arg Ile Arg Ile Arg Arg Thr
 290 295 300
 Gly Pro Ala Gln Trp Ser Gly Thr Cys Gly
 305 310

<210> 234
 <211> 93
 <212> PRT
 <213> Homo sapiens

<400> 234
 Met Arg Pro Gln Gly Pro Ala Ala Ser Pro Gln Arg Leu Arg Gly Leu
 1 5 10 15
 Leu Leu Leu Leu Leu Leu Gln Leu Pro Ala Pro Ser Ser Ala Ser Glu
 20 25 30
 Ile Pro Lys Gly Lys Gln Lys Ala His Ser Gly Arg Gly Arg Trp Trp
 35 40 45
 Thr Cys Ile Met Glu Cys Ala Tyr Lys Gly Gln Gln Glu Cys Leu Val
 50 55 60
 Glu Thr Gly Ala Leu Gly Pro Met Ala Phe Arg Val His Leu Gly Ser
 65 70 75 80
 Gln Val Gly Met Asp Ser Lys Glu Lys Arg Gly Asn Val
 85 90

<210> 235
 <211> 73
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (73)
 <223> Xaa equals stop translation
 <400> 235

Met Gly Ser Ala Ala Leu Glu Ile Leu Gly Leu Val Leu Cys Leu Val
 1 5 10 15
 Gly Trp Gly Gly Leu Ile Leu Ala Cys Gly Leu Pro Met Trp Gln Val
 20 25 30
 Thr Ala Phe Leu Asp His Asn Ile Val Thr Ala Gln Thr Thr Trp Lys
 35 40 45
 Gly Leu Trp Met Ser Cys Val Val Gln Ser Thr Gly Thr Cys Ser Ala
 50 55 60
 Lys Cys Thr Thr Arg Cys Trp Leu Xaa
 65 70

<210> 236
 <211> 349
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (283)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (293)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (325)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (326)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (349)
 <223> Xaa equals stop translation

<400> 236
 Met Leu Cys Pro Trp Arg Thr Ala Asn Leu Gly Leu Leu Ile Leu
 1 5 10 15
 Thr Ile Phe Leu Val Ala Glu Ala Glu Gly Ala Ala Gln Pro Asn Asn
 20 25 30
 Ser Leu Met Leu Gln Thr Ser Lys Glu Asn His Ala Leu Ala Ser Ser
 35 40 45
 Ser Leu Cys Met Asp Glu Lys Gln Ile Thr Gln Asn Tyr Ser Lys Val
 50 55 60

Leu Ala Glu Val Asn Thr Ser Trp Pro Val Lys Met Ala Thr Asn Ala
 65 70 75 80
 Val Leu Cys Cys Pro Pro Ile Ala Leu Arg Asn Leu Ile Ile Ile Thr
 85 90 95
 Trp Glu Ile Ile Leu Arg Gly Gln Pro Ser Cys Thr Lys Ala Tyr Lys
 100 105 110
 Lys Glu Thr Asn Glu Thr Lys Glu Thr Asn Cys Thr Asp Glu Arg Ile
 115 120 125
 Thr Trp Val Ser Arg Pro Asp Gln Asn Ser Asp Leu Gln Ile Arg Thr
 130 135 140
 Val Ala Ile Thr His Asp Gly Tyr Tyr Arg Cys Ile Met Val Thr Pro
 145 150 155 160
 Asp Gly Asn Phe His Arg Gly Tyr His Leu Gln Val Leu Val Thr Pro
 165 170 175
 Glu Val Thr Leu Phe Gln Asn Arg Asn Arg Thr Ala Val Cys Lys Ala
 180 185 190
 Val Ala Gly Lys Pro Ala Ala His Ile Ser Trp Ile Pro Glu Gly Asp
 195 200 205
 Cys Ala Thr Lys Cln Glu Tyr Trp Ser Asn Gly Thr Val Thr Val Lys
 210 215 220
 Ser Thr Cys His Trp Glu Val His Asn Val Ser Thr Val Asn Cys His
 225 230 235 240
 Val Ser His Leu Thr Gly Asn Lys Ser Leu Tyr Ile Glu Leu Leu Pro
 245 250 255
 Val Pro Gly Ala Lys Lys Ser Ser Lys Leu Tyr Ile Pro Tyr Ile Ile
 260 265 270
 Leu Thr Ile Ile Ile Leu Thr Ile Val Gly Xaa Ile Trp Leu Leu Lys
 275 280 285
 Val Asn Gly Cys Xaa Lys Tyr Lys Leu Asn Lys Pro Glu Ser Thr Pro
 290 295 300
 Val Val Glu Glu Asp Glu Met Gln Pro Tyr Ala Phe Tyr Thr Glu Lys
 305 310 315 320
 Asn Asn Pro Leu Xaa Xaa Thr Thr Asn Lys Val Lys Ala Ser Glu Ala
 325 330 335
 Leu Gln Ser Glu Val Asp Thr Asp Leu His Thr Leu Xaa
 340 345

<210> 237

<211> 17

<212> PRT

<213> Homo sapiens

<400> 237

Leu Ala Leu Tyr Ser Ala Leu Phe Ser Tyr Ser Gly Trp Asp Thr Leu
1 5 10 15

Asn

<210> 238

<211> 14

<212> PRT

<213> Homo sapiens

<400> 238

Val Thr Glu Glu Ile Lys Asn Pro Glu Arg Asn Leu Pro Leu
1 5 10

<210> 239

<211> 9

<212> PRT

<213> Homo sapiens

<400> 239

Ile Gly Ile Ser Met Pro Ile Val Thr
1 5

<210> 240

<211> 13

<212> PRT

<213> Homo sapiens

<400> 240

Ile Tyr Ile Leu Thr Asn Val Ala Tyr Tyr Thr Val Leu
1 5 10

<210> 241

<211> 11

<212> PRT

<213> Homo sapiens

<400> 241

Ser Asp Ala Val Ala Val Thr Phe Ala Asp Gln
1 5 10

<210> 242

<211> 13

<212> PRT

<213> Homo sapiens

<400> 242

Val Ala Leu Ser Cys Phe Gly Gly Leu Asn Ala Ser Ile
1 5 10

<210> 243
<211> 15
<212> PRT
<213> Homo sapiens

<400> 243
Ser Arg Leu Phe Phe Val Gly Ser Arg Glu Gly His Leu Pro Asp
1 5 10 15

<210> 244
<211> 11
<212> PRT
<213> Homo sapiens

<400> 244
Ser Phe Ser Tyr Trp Phe Phe Val Gly Leu Ser
1 5 10

<210> 245
<211> 11
<212> PRT
<213> Homo sapiens

<400> 245
Val Gly Gln Leu Tyr Leu Arg Trp Lys Glu Pro
1 5 10

<210> 246
<211> 16
<212> PRT
<213> Homo sapiens

<400> 246
Arg Pro Arg Pro Leu Lys Leu Ser Val Phe Phe Pro Ile Val Phe Cys
1 5 10 15

<210> 247
<211> 9
<212> PRT
<213> Homo sapiens

<400> 247
Asp Thr Ile Asn Ser Leu Ile Gly Ile
1 5

<210> 248
<211> 44
<212> PRT
<213> Homo sapiens

<400> 248

Ala Thr Ala Leu Pro Pro Lys Ile Val Gly Ser Ala Thr Arg Tyr Leu
 1 5 10 15
 Gln Val Leu Cys Met Ser Val Ala Ala Glu Met Asp Leu Glu Asp Gly
 20 25 30
 Gly Glu Met Pro Lys Gln Arg Asp Pro Lys Ser Asn
 35 40

<210> 249
 <211> 352
 <212> PRT
 <213> Homo sapiens

<400> 249
 Leu Leu Ala Ala Ala Cys Ile Cys Leu Leu Thr Phe Ile Asn Cys Ala
 1 5 10 15
 Tyr Val Lys Trp Gly Thr Leu Val Gln Asp Ile Phe Thr Tyr Ala Lys
 20 25 30
 Val Leu Ala Leu Ile Ala Val Ile Val Ala Gly Ile Val Arg Leu Gly
 35 40 45
 Gln Gly Ala Ser Thr His Phe Glu Asn Ser Phe Glu Gly Ser Ser Phe
 50 55 60
 Ala Val Gly Asp Ile Ala Leu Ala Leu Tyr Ser Ala Leu Phe Ser Tyr
 65 70 75 80
 Ser Gly Trp Asp Thr Leu Asn Tyr Val Thr Glu Glu Ile Lys Asn Pro
 85 90 95
 Glu Arg Asn Leu Pro Leu Ser Ile Gly Ile Ser Met Pro Ile Val Thr
 100 105 110
 Ile Ile Tyr Ile Leu Thr Asn Val Ala Tyr Tyr Thr Val Leu Asp Met
 115 120 125
 Arg Asp Ile Leu Ala Ser Asp Ala Val Ala Val Thr Phe Ala Asp Gln
 130 135 140
 Ile Phe Gly Ile Phe Asn Trp Ile Ile Pro Leu Ser Val Ala Leu Ser
 145 150 155 160
 Cys Phe Gly Gly Leu Asn Ala Ser Ile Val Ala Ala Ser Arg Leu Phe
 165 170 175
 Phe Val Gly Ser Arg Glu Gly His Leu Pro Asp Ala Ile Cys Met Ile
 180 185 190
 His Val Glu Arg Phe Thr Pro Val Pro Ser Leu Leu Phe Asn Gly Ile
 195 200 205
 Met Ala Leu Ile Tyr Leu Cys Val Glu Asp Ile Phe Gln Leu Ile Asn
 210 215 220
 Tyr Tyr Ser Phe Ser Tyr Trp Phe Phe Val Gly Leu Ser Ile Val Gly

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225          230          235          240
Gln Leu Tyr Leu Arg Trp Lys Glu Pro Asp Arg Pro Arg Pro Leu Lys
          245          250          255
Leu Ser Val Phe Phe Pro Ile Val Phe Cys Leu Cys Thr Ile Phe Leu
          260          265          270
Val Ala Val Pro Leu Tyr Ser Asp Thr Ile Asn Ser Leu Ile Gly Ile
          275          280          285
Ala Ile Ala Leu Ser Gly Leu Pro Phe Tyr Phe Leu Ile Ile Arg Val
          290          295          300
Pro Glu His Lys Arg Pro Leu Tyr Leu Arg Arg Ile Val Gly Ser Ala
305          310          315          320
Thr Arg Tyr Leu Gln Val Leu Cys Met Ser Val Ala Ala Glu Met Asp
          325          330          335
Leu Glu Asp Gly Gly Glu Met Pro Lys Gln Arg Asp Pro Lys Ser Asn
          340          345          350

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<210> 250
<211> 119
<212> PRT
<213> Homo sapiens

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<400> 250
Ala Ala Arg Gly Ser Gly Val Arg Asp Pro Leu Glu Glu Ala Val Cys
 1          5          10          15
Pro Phe Ser Asp Leu Gln Leu His Ala Gly Arg Thr Thr Ala Leu Phe
          20          25          30
Lys Ala Val Arg Gln Gly His Leu Ser Leu Gln Arg Leu Leu Ser
          35          40          45
Phe Val Cys Leu Cys Pro Ala Pro Arg Gly Gly Ala Tyr Arg Gly Arg
          50          55          60
Gln Ala Ser Leu Ser Cys Gly Gly Leu His Pro Val Arg Ala Ser Arg
          65          70          75          80
Leu Leu Cys Leu Pro Lys Gln Ala Trp Ala Met Ala Gly Ala Pro Pro
          85          90          95
Pro Val Ser Leu Pro Pro Cys Ser Leu Ile Ser Asp Cys Cys Ala Ser
          100          105          110
Asn Gln Arg Asp Ser Val Gly
          115

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<210> 251

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<211> 356
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (37)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (280)
 <223> Xaa equals any of the naturally occurring L-amino acids

<400> 251
 Leu Ser Lys Ala Phe Leu Asp Ser Pro Asn Arg Leu Leu Ala Val Glu
 1 5 10 15
 Met Asn Thr Asp His Leu Arg Leu Thr Val Pro Asn Gly Ile Gly Ala
 20 25 30
 Leu Lys Leu Arg Xaa Met Glu His Tyr Phe Ser Gln Gly Leu Ser Val
 35 40 45
 Gln Leu Phe Asn Asp Gly Ser Lys Gly Lys Leu Asn His Leu Cys Gly
 50 55 60
 Ala Asp Phe Val Lys Ser His Gln Lys Pro Pro Gln Gly Met Glu Ile
 65 70 75 80
 Lys Ser Asn Glu Arg Cys Cys Ser Phe Asp Gly Asp Ala Asp Arg Ile
 85 90 95
 Val Tyr Tyr Tyr His Asp Ala Asp Gly His Phe His Leu Ile Asp Gly
 100 105 110
 Asp Lys Ile Ala Thr Leu Ile Ser Ser Phe Leu Lys Glu Leu Leu Val
 115 120 125
 Glu Ile Gly Glu Ser Leu Asn Ile Gly Val Val Gln Thr Ala Tyr Ala
 130 135 140
 Asn Gly Ser Ser Thr Arg Tyr Leu Glu Glu Val Met Lys Val Pro Val
 145 150 155 160
 Tyr Cys Thr Lys Thr Gly Val Lys His Leu His His Lys Ala Gln Glu
 165 170 175
 Phe Asp Ile Gly Val Tyr Phe Glu Ala Asn Gly His Gly Thr Ala Leu
 180 185 190
 Phe Ser Thr Ala Val Glu Met Lys Ile Lys Gln Ser Ala Glu Gln Leu
 195 200 205
 Glu Asp Lys Lys Arg Lys Ala Ala Lys Met Leu Glu Asn Ile Ile Asp
 210 215 220
 Leu Phe Asn Gln Ala Ala Gly Asp Ala Ile Ser Asp Met Leu Val Ile
 225 230 235 240

Glu Ala Ile Leu Ala Leu Lys Gly Leu Thr Val Gln Gln Trp Asp Ala
 245 250 255
 Leu Tyr Thr Asp Leu Pro Asn Arg Gln Leu Lys Val Gln Val Ala Asp
 260 265 270
 Arg Arg Val Ile Ser Thr Thr Xaa Ala Glu Arg Gln Ala Val Thr Pro
 275 280 285
 Pro Gly Leu Gln Glu Ala Ile Asn Asp Leu Val Lys Lys Tyr Lys Leu
 290 295 300
 Ser Arg Ala Phe Val Arg Pro Ser Gly Thr Glu Asp Val Val Arg Val
 305 310 315 320
 Tyr Ala Glu Ala Asp Ser Gln Glu Ser Ala Asp His Leu Ala His Glu
 325 330 335
 Val Ser Leu Ala Val Phe Gln Leu Ala Gly Gly Ile Gly Glu Arg Pro
 340 345 350
 Gln Pro Gly Phe
 355

<210> 252
 <211> 26
 <212> PRT
 <213> Homo sapiens

<400> 252
 Leu Ser Lys Ala Phe Leu Asp Ser Pro Asn Arg Leu Leu Ala Val Glu
 1 5 10 15
 Met Asn Thr Asp His Leu Arg Leu Thr Val
 20 25

<210> 253
 <211> 28
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (11)
 <223> Xaa equals any of the naturally occurring L-amino acids

<400> 253
 Pro Asn Gly Ile Gly Ala Leu Lys Leu Arg Xaa Met Glu His Tyr Phe
 1 5 10 15
 Ser Gln Gly Leu Ser Val Gln Leu Phe Asn Asp Gly
 20 25

<210> 254
 <211> 28

<212> PRT

<213> Homo sapiens

<400> 254

Ser Lys Gly Lys Leu Asn His Leu Cys Gly Ala Asp Phe Val Lys Ser
1 5 10 15

His Gln Lys Pro Pro Gln Gly Met Glu Ile Lys Ser
20 25

<210> 255

<211> 28

<212> PRT

<213> Homo sapiens

<400> 255

Asn Glu Arg Cys Cys Ser Phe Asp Gly Asp Ala Asp Arg Ile Val Tyr
1 5 10 15

Tyr Tyr His Asp Ala Asp Gly His Phe His Leu Ile
20 25

<210> 256

<211> 28

<212> PRT

<213> Homo sapiens

<400> 256

Asp Gly Asp Lys Ile Ala Thr Leu Ile Ser Ser Phe Leu Lys Glu Leu
1 5 10 15

Leu Val Glu Ile Gly Glu Ser Leu Asn Ile Gly Val
20 25

<210> 257

<211> 28

<212> PRT

<213> Homo sapiens

<400> 257

Val Gln Thr Ala Tyr Ala Asn Gly Ser Ser Thr Arg Tyr Leu Glu Glu
1 5 10 15

Val Met Lys Val Pro Val Tyr Cys Thr Lys Thr Gly
20 25

<210> 258

<211> 28

<212> PRT

<213> Homo sapiens

<400> 258

Val Lys His Leu His His Lys Ala Gln Glu Phe Asp Ile Gly Val Tyr
1 5 10 15

Phe Glu Ala Asn Gly His Gly Thr Ala Leu Phe Ser
20 25

<210> 259
<211> 28
<212> PRT
<213> Homo sapiens

<400> 259
Thr Ala Val Glu Met Lys Ile Lys Gln Ser Ala Glu Gln Leu Glu Asp
1 5 10 15
Lys Lys Arg Lys Ala Ala Lys Met Leu Glu Asn Ile
20 25

<210> 260
<211> 28
<212> PRT
<213> Homo sapiens

<400> 260
Ile Asp Leu Phe Asn Gln Ala Ala Gly Asp Ala Ile Ser Asp Met Leu
1 5 10 15
Val Ile Glu Ala Ile Leu Ala Leu Lys Gly Leu Thr
20 25

<210> 261
<211> 28
<212> PRT
<213> Homo sapiens

<400> 261
Val Gln Gln Trp Asp Ala Leu Tyr Thr Asp Leu Pro Asn Arg Gln Leu
1 5 10 15
Lys Val Gln Val Ala Asp Arg Arg Val Ile Ser Thr
20 25

<210> 262
<211> 28
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (2)
<223> Xaa equals any of the naturally occurring L-amino acids

<400> 262
Thr Xaa Ala Glu Arg Gln Ala Val Thr Pro Pro Gly Leu Gln Glu Ala
1 5 10 15
Ile Asn Asp Leu Val Lys Lys Tyr Lys Leu Ser Arg
20 25

<210> 263

<211> 24

<212> PRT

<213> Homo sapiens

<400> 263

Ala Phe Val Arg Pro Ser Gly Thr Glu Asp Val Val Arg Val Tyr Ala
1 5 10 15

Glu Ala Asp Ser Gln Glu Ser Ala
20

<210> 264

<211> 26

<212> PRT

<213> Homo sapiens

<400> 264

Asp His Leu Ala His Glu Val Ser Leu Ala Val Phe Gln Leu Ala Gly
1 5 10 15

Gly Ile Gly Glu Arg Pro Gln Pro Gly Phe
20 25

<210> 265

<211> 443

<212> PRT

<213> Homo sapiens

<400> 265

Gly Thr Arg Ala Ala Pro Gly Leu Gly Ala Trp Gly Arg Arg Ser Pro
1 5 10 15

Pro Ser Phe Ser Pro Pro Arg Pro Arg Arg Pro Gly Val Met Ala Gly
20 25 30

Leu Asn Cys Gly Val Ser Ile Ala Leu Leu Gly Val Leu Leu Leu Gly
35 40 45

Ala Ala Arg Leu Pro Arg Gly Ala Glu Ala Phe Glu Ile Ala Leu Pro
50 55 60

Arg Glu Ser Asn Ile Thr Val Leu Ile Lys Leu Gly Thr Pro Thr Leu
65 70 75 80

Leu Ala Lys Pro Cys Tyr Ile Val Ile Ser Lys Arg His Ile Thr Met
85 90 95

Leu Ser Ile Lys Ser Gly Glu Arg Ile Val Phe Thr Phe Ser Cys Gln
100 105 110

Ser Pro Glu Asn His Phe Val Ile Glu Ile Gln Lys Asn Ile Asp Cys
115 120 125

Met Ser Gly Pro Cys Pro Phe Gly Glu Val Gln Leu Gln Pro Ser Thr

130	135	140
Ser Leu Leu Pro Thr	Leu Asn Arg Thr Phe	Ile Trp Asp Val Lys Ala
145	150	155 160
His Lys Ser Ile Gly	Leu Glu Leu Gln Phe	Ser Ile Pro Arg Leu Arg
165	170	175
Gln Ile Gly Pro Gly	Glu Ser Cys Pro Asp	Gly Val Thr His Ser Ile
180	185	190
Ser Gly Arg Ile Asp	Ala Thr Val Val Arg	Ile Gly Thr Phe Cys Ser
195	200	205
Asn Gly Thr Val Ser	Arg Ile Lys Met Gln	Glu Gly Val Lys Met Ala
210	215	220
Leu His Leu Pro Trp	Phe His Pro Arg Asn	Val Ser Gly Phe Ser Ile
225	230	235 240
Ala Asn Arg Ser Ser	Ile Lys Arg Leu Cys	Ile Ile Glu Ser Val Phe
245	250	255
Glu Gly Glu Gly Ser	Ala Thr Leu Met Ser	Ala Asn Tyr Pro Glu Gly
260	265	270
Phe Pro Glu Asp Glu	Leu Met Thr Trp Gln	Phe Val Val Pro Ala His
275	280	285
Leu Arg Ala Ser Val	Ser Phe Leu Asn Phe	Asn Leu Ser Asn Cys Glu
290	295	300
Arg Lys Glu Glu Arg	Val Glu Tyr Tyr Ile	Pro Gly Ser Thr Thr Asn
305	310	315 320
Pro Glu Val Phe Lys	Leu Glu Asp Lys Gln	Pro Gly Asn Met Ala Gly
325	330	335
Asn Phe Asn Leu Ser	Leu Gln Gly Cys Asp	Gln Asp Ala Gln Ser Pro
340	345	350
Gly Ile Leu Arg Leu	Gln Phe Gln Val Leu	Val Gln His Pro Gln Asn
355	360	365
Glu Ser Asn Lys Ile	Tyr Val Val Asp Leu	Ser Asn Glu Arg Ala Met
370	375	380
Ser Leu Thr Ile Glu	Pro Arg Pro Val Lys	Gln Ser Arg Lys Phe Val
385	390	395 400
Pro Gly Cys Phe Val	Cys Leu Glu Ser Arg	Thr Cys Ser Ser Asn Leu
405	410	415
Thr Leu Thr Ser Gly	Ser Lys His Lys Ile	Ser Phe Leu Cys Asp Asp
420	425	430
Leu Thr Arg Leu Trp	Met Asn Val Glu Lys	Pro
435	440	

<210> 266
<211> 159
<212> PRT
<213> Homo sapiens

<400> 266
Phe Glu Ile Ala Leu Pro Arg Glu Ser Asn Ile Thr Val Leu Ile Lys
1 5 10 15
Leu Gly Thr Pro Thr Leu Leu Ala Lys Pro Cys Tyr Ile Val Ile Ser
20 25 30
Lys Arg His Ile Thr Met Leu Ser Ile Lys Ser Gly Glu Arg Ile Val
35 40 45
Phe Thr Phe Ser Cys Gln Ser Pro Glu Asn His Phe Val Ile Glu Ile
50 55 60
Gln Lys Asn Ile Asp Cys Met Ser Gly Pro Cys Pro Phe Gly Glu Val
65 70 75 80
Gln Leu Gln Pro Ser Thr Ser Leu Leu Pro Thr Leu Asn Arg Thr Phe
85 90 95
Ile Trp Asp Val Lys Ala His Lys Ser Ile Gly Leu Glu Leu Gln Phe
100 105 110
Ser Ile Pro Arg Leu Arg Gln Ile Gly Pro Gly Glu Ser Cys Pro Asp
115 120 125
Gly Val Thr His Ser Ile Ser Gly Arg Ile Asp Ala Thr Val Val Arg
130 135 140
Ile Gly Thr Phe Cys Ser Asn Gly Thr Val Ser Arg Ile Lys Met
145 150 155

<210> 267
<211> 9
<212> PRT
<213> Homo sapiens

<400> 267
Phe Val Arg Asp Pro Phe Val Arg Leu
1 5

<210> 268
<211> 13
<212> PRT
<213> Homo sapiens

<400> 268
Phe Leu Phe Val Arg Asp Pro Phe Val Arg Leu Ile Ser
1 5 10

<210> 269

<211> 15
 <212> PRT
 <213> Homo sapiens

<400> 269

Phe Leu Phe Val Arg Asp Pro Phe Val Arg Leu Ile Ser Ala Phe
 1 5 10 15

<210> 270
 <211> 380
 <212> PRT
 <213> Homo sapiens

<400> 270

Tyr Leu His Thr Ser Phe Ser Arg Pro His Thr Gly Pro Pro Leu Pro
 1 5 10 15
 Thr Pro Gly Pro Asp Arg Asp Arg Glu Leu Thr Ala Asp Ser Asp Val
 20 25 30
 Asp Glu Phe Leu Asp Lys Phe Leu Ser Ala Gly Val Lys Gln Ser Asp
 35 40 45
 Leu Pro Arg Lys Glu Thr Glu Gln Pro Pro Ala Pro Gly Ser Met Glu
 50 55 60
 Glu Asn Val Arg Gly Tyr Asp Trp Ser Pro Arg Asp Ala Arg Arg Ser
 65 70 75 80
 Pro Asp Gln Gly Arg Gln Gln Ala Glu Arg Arg Ser Val Leu Arg Gly
 85 90 95
 Phe Cys Ala Asn Ser Ser Leu Ala Phe Pro Thr Lys Glu Arg Ala Phe
 100 105 110
 Asp Asp Ile Pro Asn Ser Glu Leu Ser His Leu Ile Val Asp Asp Arg
 115 120 125
 His Gly Ala Ile Tyr Cys Tyr Val Pro Lys Val Ala Cys Thr Asn Trp
 130 135 140
 Lys Arg Val Met Ile Val Leu Ser Gly Ser Leu Leu His Arg Gly Ala
 145 150 155 160
 Pro Tyr Arg Asp Pro Leu Arg Ile Pro Arg Glu His Val His Asn Ala
 165 170 175
 Ser Ala His Leu Thr Phe Asn Lys Phe Trp Arg Arg Tyr Gly Lys Leu
 180 185 190
 Ser Arg His Leu Met Lys Val Lys Leu Lys Lys Tyr Thr Lys Phe Leu
 195 200 205
 Phe Val Arg Asp Pro Phe Val Arg Leu Ile Ser Ala Phe Arg Ser Lys
 210 215 220
 Phe Glu Leu Glu Asn Glu Glu Phe Tyr Arg Lys Phe Ala Val Pro Met
 225 230 235 240

Leu Arg Leu Tyr Ala Asn His Thr Ser Leu Pro Ala Ser Ala Arg Glu
 245 250 255
 Ala Phe Arg Ala Gly Leu Lys Val Ser Phe Ala Asn Phe Ile Gln Tyr
 260 265 270
 Leu Leu Asp Pro His Thr Glu Lys Leu Ala Pro Phe Asn Glu His Trp
 275 280 285
 Arg Gln Val Tyr Arg Leu Cys His Pro Cys Gln Ile Asp Tyr Asp Phe
 290 295 300
 Val Gly Lys Leu Glu Thr Leu Asp Glu Asp Ala Ala Gln Leu Leu Gln
 305 310 315 320
 Leu Leu Gln Val Asp Arg Gln Leu Arg Phe Pro Pro Ser Tyr Arg Asn
 325 330 335
 Arg Thr Ala Ser Ser Trp Glu Glu Asp Trp Phe Ala Lys Ile Pro Leu
 340 345 350
 Ala Trp Arg Gln Gln Leu Tyr Lys Leu Tyr Glu Ala Asp Phe Val Leu
 355 360 365
 Phe Gly Tyr Pro Lys Pro Glu Asn Leu Leu Arg Asp
 370 375 380

 <210> 271
 <211> 274
 <212> PRT
 <213> Homo sapiens

 <400> 271
 Lys Leu Val Arg Leu Gln Val Pro Val Arg Asn Ser Arg Val Asp Pro
 1 5 10 15
 Arg Val Arg Ser Lys Ile Gly Ser Arg Arg Trp Met Leu Gln Leu Ile
 20 25 30
 Met Gln Leu Gly Ser Val Leu Leu Thr Arg Cys Pro Phe Trp Gly Cys
 35 40 45
 Phe Ser Gln Leu Met Leu Tyr Ala Glu Arg Ala Glu Ala Arg Arg Lys
 50 55 60
 Pro Asp Ile Pro Val Pro Tyr Leu Tyr Phe Asp Met Gly Ala Ala Val
 65 70 75 80
 Leu Cys Ala Ser Phe Met Ser Phe Gly Val Lys Arg Arg Trp Phe Ala
 85 90 95
 Leu Gly Ala Ala Leu Gln Leu Ala Ile Ser Thr Tyr Ala Ala Tyr Ile
 100 105 110
 Gly Gly Tyr Val His Tyr Gly Asp Trp Leu Lys Val Arg Met Tyr Ser
 115 120 125

Arg Thr Val Ala Ile Ile Gly Gly Phe Leu Val Leu Ala Ser Gly Ala
 130 135 140
 Gly Glu Leu Tyr Arg Arg Lys Pro Arg Ser Arg Ser Leu Gln Ser Thr
 145 150 155 160
 Gly Gln Val Phe Leu Gly Ile Tyr Leu Ile Cys Val Ala Tyr Ser Leu
 165 170 175
 Gln His Ser Lys Glu Asp Arg Leu Ala Tyr Leu Asn His Leu Pro Gly
 180 185 190
 Gly Glu Leu Met Ile Gln Leu Phe Phe Val Leu Tyr Gly Ile Leu Ala
 195 200 205
 Leu Ala Phe Leu Ser Gly Tyr Tyr Val Thr Leu Ala Ala Gln Ile Leu
 210 215 220
 Ala Val Leu Leu Pro Pro Val Met Leu Leu Ile Asp Gly Asn Val Ala
 225 230 235 240
 Tyr Trp His Asn Thr Arg Arg Val Glu Phe Trp Asn Gln Met Lys Leu
 245 250 255
 Leu Gly Glu Ser Val Gly Ile Phe Gly Thr Ala Val Ile Leu Ala Thr
 260 265 270

Asp Gly

<210> 272

<211> 203

<212> PRT

<213> Homo sapiens

<400> 272

Met Gln Leu Gly Ser Val Leu Leu Thr Arg Cys Pro Phe Trp Gly Cys
 1 5 10 15
 Phe Ser Gln Leu Met Leu Tyr Ala Glu Arg Ala Glu Ala Arg Arg Lys
 20 25 30
 Pro Asp Ile Pro Val Pro Tyr Leu Tyr Phe Asp Met Gly Ala Ala Val
 35 40 45
 Leu Cys Ala Ser Phe Met Ser Phe Gly Val Lys Arg Arg Trp Phe Ala
 50 55 60
 Leu Gly Ala Ala Leu Gln Leu Ala Ile Ser Thr Tyr Ala Ala Tyr Ile
 65 70 75 80
 Gly Gly Tyr Val His Tyr Gly Asp Trp Leu Lys Val Arg Met Tyr Ser
 85 90 95
 Arg Thr Val Ala Ile Ile Gly Gly Phe Leu Val Leu Ala Ser Gly Ala
 100 105 110
 Gly Glu Leu Tyr Arg Arg Lys Pro Arg Ser Arg Ser Leu Gln Ser Thr

115 120 125
 Gly Gln Val Phe Leu Gly Ile Tyr Leu Ile Cys Val Ala Tyr Ser Leu
 130 135 140
 Gln His Ser Lys Glu Asp Arg Leu Ala Tyr Leu Asn His Leu Pro Gly
 145 150 155 160
 Gly Glu Leu Met Ile Gln Leu Phe Phe Val Leu Tyr Gly Ile Leu Ala
 165 170 175
 Pro Gly Leu Ser Val Arg Leu Leu Arg Asp Pro Arg Cys Pro Asp Pro
 180 185 190
 Gly Cys Thr Ala Ala Pro Cys His Ala Ala His
 195 200

 <210> 273
 <211> 407
 <212> PRT
 <213> Homo sapiens

 <400> 273
 Ser Asn Glu Ile Leu Leu Ser Phe Pro Gln Asn Tyr Tyr Ile Gln Trp
 1 5 10 15
 Leu Asn Gly Ser Leu Ile His Gly Leu Trp Asn Leu Ala Ser Leu Phe
 20 25 30
 Ser Asn Leu Cys Leu Phe Val Leu Met Pro Phe Ala Phe Phe Phe Leu
 35 40 45
 Glu Ser Glu Gly Phe Ala Gly Leu Lys Lys Gly Ile Arg Ala Arg Ile
 50 55 60
 Leu Glu Thr Leu Val Met Leu Leu Leu Ala Leu Leu Ile Leu Gly
 65 70 75 80
 Ile Val Trp Val Ala Ser Ala Leu Ile Asp Asn Asp Ala Ala Ser Met
 85 90 95
 Glu Ser Leu Tyr Asp Leu Trp Glu Phe Tyr Leu Pro Tyr Leu Tyr Ser
 100 105 110
 Cys Ile Ser Leu Met Gly Cys Leu Leu Leu Leu Cys Thr Pro Val
 115 120 125
 Gly Leu Ser Arg Met Phe Thr Val Met Gly His Leu Leu Val Lys Pro
 130 135 140
 Thr Ile Leu Glu Asp Leu Asp Glu Gln Ile Tyr Ile Ile Thr Leu Glu
 145 150 155 160
 Glu Glu Ala Leu Gln Arg Arg Leu Asn Gly Leu Ser Ser Ser Val Glu
 165 170 175
 Tyr Asn Ile Met Glu Leu Glu Gln Glu Leu Glu Asn Val Lys Thr Leu
 180 185 190

Lys Thr Lys Leu Glu Arg Arg Lys Lys Ala Ser Ala Trp Glu Arg Asn
 195 200 205
 Leu Val Tyr Pro Ala Val Met Val Leu Leu Leu Ile Glu Thr Ser Ile
 210 215 220
 Ser Val Leu Leu Val Ala Cys Asn Ile Leu Cys Leu Leu Val Asp Glu
 225 230 235 240
 Thr Ala Met Pro Lys Gly Thr Arg Gly Pro Gly Ile Gly Asn Ala Ser
 245 250 255
 Leu Ser Thr Phe Gly Phe Val Gly Ala Ala Leu Glu Ile Ile Leu Ile
 260 265 270
 Phe Tyr Leu Met Val Ser Ser Val Val Gly Phe Tyr Ser Leu Arg Phe
 275 280 285
 Phe Gly Asn Phe Thr Pro Lys Lys Asp Asp Thr Thr Met Thr Lys Ile
 290 295 300
 Ile Gly Asn Cys Val Ser Ile Leu Val Leu Ser Ser Ala Leu Pro Val
 305 310 315 320
 Met Ser Arg Thr Leu Gly Ile Thr Arg Phe Asp Leu Leu Gly Asp Phe
 325 330 335
 Gly Arg Phe Asn Trp Leu Gly Asn Phe Tyr Ile Val Leu Ser Tyr Asn
 340 345 350
 Leu Leu Phe Ala Ile Val Thr Thr Leu Cys Leu Val Arg Lys Phe Thr
 355 360 365
 Ser Ala Val Arg Glu Glu Leu Phe Lys Ala Leu Gly Leu His Lys Leu
 370 375 380
 His Leu Pro Asn Thr Ser Arg Asp Ser Glu Thr Ala Lys Pro Ser Val
 385 390 395 400
 Asn Gly His Gln Lys Ala Leu
 405

<210> 274
 <211> 165
 <212> PRT
 <213> Homo sapiens

<400> 274
 Arg Ser Tyr Met Gln Ser Val Trp Thr Glu Glu Ser Gln Cys Thr Leu
 1 5 10 15
 Leu Asn Ala Ser Ile Thr Glu Thr Phe Asn Cys Ser Phe Ser Cys Gly
 20 25 30
 Pro Asp Cys Trp Lys Leu Ser Gln Tyr Pro Cys Leu Gln Val Tyr Val
 35 40 45

Asn Leu Thr Ser Ser Gly Glu Lys Leu Leu Leu Tyr His Thr Glu Glu
 50 55 60
 Thr Ile Lys Ile Asn Gln Lys Cys Ser Tyr Ile Pro Lys Cys Gly Lys
 65 70 75 80
 Asn Phe Glu Glu Ser Met Ser Leu Val Asn Val Val Met Glu Asn Phe
 85 90 95
 Arg Lys Tyr Gln His Phe Ser Cys Tyr Ser Asp Pro Glu Gly Asn Gln
 100 105 110
 Lys Ser Val Ile Leu Thr Lys Leu Tyr Ser Ser Asn Val Leu Phe His
 115 120 125
 Ser Leu Phe Trp Pro Thr Cys Met Met Ala Gly Gly Val Ala Ile Val
 130 135 140
 Ala Met Val Lys Leu Thr Gln Tyr Leu Ser Leu Leu Cys Glu Arg Ile
 145 150 155 160
 Gln Arg Ile Asn Arg
 165

<210> 275
 <211> 155
 <212> PRT
 <213> Homo sapiens

<400> 275
 Ala Phe Ala His Leu Gln Leu Gly Pro Met Trp Lys Leu Trp Arg Ala
 1 5 10 15
 Glu Glu Gly Ala Ala Ala Leu Gly Gly Ala Leu Phe Leu Leu Phe
 20 25 30
 Ala Leu Gly Val Arg Gln Leu Leu Lys Gln Arg Arg Pro Met Gly Phe
 35 40 45
 Pro Pro Gly Pro Pro Gly Leu Pro Phe Ile Gly Asn Ile Tyr Ser Leu
 50 55 60
 Ala Ala Ser Ser Glu Leu Pro His Val Tyr Met Arg Lys Gln Ser Gln
 65 70 75 80
 Val Tyr Gly Glu Val Gln Pro Arg Arg Ala Pro Gly Arg Glu Gly Arg
 85 90 95
 Gln Ala Gly Pro Gly Trp Pro Gly Pro Ser Trp Leu Asp Leu Trp Pro
 100 105 110
 Pro Leu Gly Arg Leu Val Gly Thr Ser Pro Cys Ala Gly Cys Pro Leu
 115 120 125
 Arg Asp Thr Arg Phe Pro Gly Leu Glu Gly Arg Ser Pro Arg Arg Arg
 130 135 140
 Ala Pro Leu Gln Gly Glu Pro Arg Pro Cys Arg

145

150

155

<210> 276

<211> 42

<212> PRT

<213> Homo sapiens

<400> 276

Met Arg Val Arg Ile Gly Leu Thr Leu Leu Leu Cys Ala Val Leu Leu
1 5 10 15
Ser Leu Ala Ser Ala Ser Ser Asp Glu Glu Gly Ser Gln Asp Glu Ser
20 25 30
Leu Gly Phe Gln Asp Tyr Phe Asp Ile Arg
35 40

<210> 277

<211> 155

<212> PRT

<213> Homo sapiens

<400> 277

Met Ala Arg Gly Ser Leu Arg Arg Leu Leu Arg Leu Leu Val Leu Gly
1 5 10 15
Leu Trp Leu Ala Leu Leu Arg Ser Val Ala Gly Glu Gln Ala Pro Gly
20 25 30
Thr Ala Pro Cys Ser Arg Gly Ser Ser Trp Ser Ala Asp Leu Asp Lys
35 40 45
Cys Met Asp Cys Ser Thr Ser Cys Pro Leu Pro Ala Ala Leu Ala His
50 55 60
Pro Trp Gly Arg Ser Glu Pro Asp Leu Arg Ala Gly Ala Ala Phe Trp
65 70 75 80
Leu Phe Gly Leu Glu Thr Met Pro Gln Glu Arg Glu Val His His Pro
85 90 95
His Arg Gly Asp Arg Arg Arg Gly Leu Pro Ser Cys Gly Ala Asp Pro
100 105 110
Val Thr Met Cys Pro Leu Pro Ala Gly Ala Arg Pro Leu Ile Ile His
115 120 125
Ser Ser Ile Leu Glu Pro Val Ser Ala Ser Gln Thr Arg Arg Glu Pro
130 135 140
Ser Ser Ser Asn His Lys Gly Gly Gly Gly Arg
145 150 155

<210> 278

<211> 207

<212> PRT

<213> Homo sapiens

<400> 278

Gly Thr Ser Phe Leu Asp Pro Thr Leu Ser Leu Phe Val Leu Glu Lys
 1 5 10 15
 Phe Asn Leu Pro Ala Gly Tyr Val Gly Leu Val Phe Leu Gly Met Ala
 20 25 30
 Leu Ser Tyr Ala Ile Ser Ser Pro Leu Phe Gly Leu Leu Ser Asp Lys
 35 40 45
 Arg Pro Pro Leu Arg Lys Trp Leu Leu Val Phe Gly Asn Leu Ile Thr
 50 55 60
 Ala Gly Cys Tyr Met Leu Leu Gly Pro Val Pro Ile Leu His Ile Lys
 65 70 75 80
 Ser Gln Leu Trp Leu Leu Val Leu Ile Leu Val Val Ser Gly Leu Ser
 85 90 95
 Ala Gly Met Ser Ile Ile Pro Thr Phe Pro Glu Ile Leu Ser Cys Ala
 100 105 110
 His Glu Asn Gly Phe Glu Glu Gly Leu Ser Thr Leu Gly Leu Val Ser
 115 120 125
 Gly Leu Phe Ser Ala Met Trp Ser Ile Gly Ala Phe Met Gly Pro Thr
 130 135 140
 Leu Gly Gly Phe Leu Tyr Glu Lys Ile Gly Phe Glu Trp Ala Ala Ala
 145 150 155 160
 Ile Gln Gly Leu Trp Ala Leu Ile Ser Gly Leu Ala Met Gly Leu Phe
 165 170 175
 Tyr Leu Leu Glu Tyr Ser Arg Arg Lys Arg Ser Lys Ser Gln Asn Ile
 180 185 190
 Leu Ser Thr Glu Glu Glu Arg Thr Thr Leu Leu Pro Asn Glu Thr
 195 200 205

<210> 279

<211> 85

<212> PRT

<213> Homo sapiens

<400> 279

Gly Thr Arg Glu Ala Arg Leu Arg Asp Leu Thr Arg Phe Tyr Asp Lys
 1 5 10 15
 Val Leu Ser Leu His Glu Asp Ser Thr Thr Pro Val Ala Asn Pro Leu
 20 25 30
 Leu Ala Phe Thr Leu Ile Lys Arg Leu Gln Ser Asp Trp Arg Asn Val
 35 40 45
 Val His Ser Leu Glu Ala Ser Glu Asn Ile Arg Ala Leu Lys Asp Gly

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      50              55              60
Tyr Glu Lys Val Glu Gln Asp Leu Pro Ala Phe Glu Asp Leu Glu Gly
 65              70              75              80
Ala Ala Arg Ala Leu
      85

<210> 280
<211> 7
<212> PRT
<213> Homo sapiens

<400> 280
Ala Leu Met Arg Leu Gln Asp
 1              5

<210> 281
<211> 7
<212> PRT
<213> Homo sapiens

<400> 281
Val Glu Ala Gly Gly Ala Thr
 1              5

<210> 282
<211> 489
<212> PRT
<213> Homo sapiens

<400> 282
Gly Thr Arg Glu Ala Arg Leu Arg Asp Leu Thr Arg Phe Tyr Asp Lys
 1              5              10              15
Val Leu Ser Leu His Glu Asp Ser Thr Thr Pro Val Ala Asn Pro Leu
      20              25              30
Leu Ala Phe Thr Leu Ile Lys Arg Leu Gln Ser Asp Trp Arg Asn Val
      35              40              45
Val His Ser Leu Glu Ala Ser Glu Asn Ile Arg Ala Leu Lys Asp Gly
      50              55              60
Tyr Glu Lys Val Glu Gln Asp Leu Pro Ala Phe Glu Asp Leu Glu Gly
      65              70              75              80
Ala Ala Arg Ala Leu Met Arg Leu Gln Asp Val Tyr Met Leu Asn Val
      85              90              95
Lys Gly Leu Ala Arg Gly Val Phe Gln Arg Val Thr Gly Ser Ala Ile
      100              105              110
Thr Asp Leu Tyr Ser Pro Lys Arg Leu Phe Ser Leu Thr Gly Asp Asp
      115              120              125

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Cys Phe Gln Val Gly Lys Val Ala Tyr Asp Met Gly Asp Tyr Tyr His
 130 135 140
 Ala Ile Pro Trp Leu Glu Glu Ala Val Ser Leu Phe Arg Gly Ser Tyr
 145 150 155 160
 Gly Glu Trp Lys Thr Glu Asp Glu Ala Ser Leu Glu Asp Ala Leu Asp
 165 170 175
 His Leu Ala Phe Ala Tyr Phe Arg Ala Gly Asn Val Ser Cys Ala Leu
 180 185 190
 Ser Leu Ser Arg Glu Phe Leu Leu Tyr Ser Pro Asp Asn Lys Arg Met
 195 200 205
 Ala Arg Asn Val Leu Lys Tyr Glu Arg Leu Leu Ala Glu Ser Pro Asn
 210 215 220
 His Val Val Ala Glu Ala Val Ile Gln Arg Pro Asn Ile Pro His Leu
 225 230 235 240
 Gln Thr Arg Asp Thr Tyr Glu Gly Leu Cys Gln Thr Leu Gly Ser Gln
 245 250 255
 Pro Thr Leu Tyr Gln Ile Pro Ser Leu Tyr Cys Ser Tyr Glu Thr Asn
 260 265 270
 Ser Asn Ala Tyr Leu Leu Leu Gln Pro Ile Arg Lys Glu Val Ile His
 275 280 285
 Leu Glu Pro Tyr Ile Ala Leu Tyr His Asp Phe Val Ser Asp Ser Glu
 290 295 300
 Ala Gln Lys Ile Arg Glu Leu Ala Glu Pro Trp Leu Gln Arg Ser Val
 305 310 315 320
 Val Ala Ser Gly Glu Lys Gln Leu Gln Val Glu Tyr Arg Ile Ser Lys
 325 330 335
 Ser Ala Trp Leu Lys Asp Thr Val Asp Leu Lys Leu Val Thr Leu Asn
 340 345 350
 His Arg Ile Ala Ala Leu Thr Gly Leu Asp Val Arg Pro Pro Tyr Ala
 355 360 365
 Glu Tyr Leu Gln Val Val Asn Tyr Gly Ile Gly Gly His Tyr Glu Pro
 370 375 380
 His Phe Asp His Ala Thr Ser Pro Ser Ser Pro Leu Tyr Arg Met Lys
 385 390 395 400
 Ser Gly Asn Arg Val Ala Thr Phe Met Ile Tyr Leu Ser Ser Val Glu
 405 410 415
 Ala Gly Gly Ala Thr Ala Phe Ile Tyr Ala Asn Leu Ser Val Pro Val
 420 425 430
 Val Arg Asn Ala Ala Leu Phe Trp Trp Asn Leu His Arg Ser Gly Glu
 435 440 445

Gly Asp Ser Asp Thr Leu His Ala Gly Cys Pro Val Leu Val Gly Asp
 450 455 460

Lys Trp Val Ala Asn Lys Trp Ile His Glu Tyr Gly Gln Glu Phe Arg
 465 470 475 480

Arg Pro Cys Ser Ser Ser Pro Glu Asp
 485

<210> 283

<211> 136

<212> PRT

<213> Homo sapiens

<400> 283

Ile Gln Pro Ser His Ala Ala Leu Leu His Cys Arg Ser Thr Phe Arg
 1 5 10 15

Lys Thr Glu Cys Leu Asp Pro Trp Trp Val Arg Arg Gln Leu Leu Gly
 20 25 30

Met Ala Gly Ile Gly Gly Leu Gln Lys Met Lys Ala Pro His Thr Gly
 35 40 45

Val Leu His Leu Gly Ser Val Trp Val Phe Leu Gly Pro Phe Leu Leu
 50 55 60

Gly Val Gly Tyr Thr Leu Thr Phe Asn Pro Leu Ser Gly Cys Met Ser
 65 70 75 80

Thr Val Arg Trp Leu Asn Ser Asn Ile Thr Ala Asn Arg Thr Leu Ser
 85 90 95

Arg Ser Val Cys His Val Thr Pro Leu His Arg Ser Leu Ser Pro His
 100 105 110

Asp Gly Glu Tyr Leu Arg Gln Met Leu Leu Asn Ser Ser Ser Arg Ala
 115 120 125

Gly Glu Ala Gly Ser Trp Gly Tyr
 130 135

<210> 284

<211> 86

<212> PRT

<213> Homo sapiens

<400> 284

Cys Ser Ser Pro Pro Gly Arg Leu Pro Trp Cys Trp Thr Ala Pro Arg
 1 5 10 15

Thr Leu Gly Lys His Gly Ser Leu Ile Ser Thr Leu Arg Leu Thr Ala
 20 25 30

Pro Leu His Leu Ala Trp Lys Met Met Leu Ser Arg Lys Ala Leu Phe
 35 40 45

Val Leu Leu Asn Thr Pro Val Leu Phe His Ala Leu Glu Gly Arg Leu
 50 55 60
 Phe Ser Lys Leu Cys His His His Thr Ile Gln Arg Thr Leu Thr Val
 65 70 75 80
 Pro Lys Phe Arg Ser Ser
 85

<210> 285
 <211> 75
 <212> PRT
 <213> Homo sapiens

<400> 285
 Arg Ser Pro Thr Ser Arg Val Gln Leu Leu Lys Arg Gln Ser Cys Pro
 1 5 10 15
 Cys Gln Arg Asn Asp Leu Asn Glu Glu Pro Gln His Phe Thr His Tyr
 20 25 30
 Ala Ile Tyr Asp Phe Ile Val Lys Gly Ser Cys Phe Cys Asn Gly His
 35 40 45
 Ala Asp Gln Cys Ile Pro Val His Gly Phe Arg Pro Val Lys Ala Pro
 50 55 60
 Gly Thr Phe His Met Val His Gly Lys Cys Met
 65 70 75

<210> 286
 <211> 296
 <212> PRT
 <213> Homo sapiens

<400> 286
 His Asn Thr Ala Gly Ser His Cys Gln His Cys Ala Pro Leu Tyr Asn
 1 5 10 15
 Asp Arg Pro Trp Glu Ala Ala Asp Gly Lys Thr Gly Ala Pro Asn Glu
 20 25 30
 Cys Arg Thr Cys Lys Cys Asn Gly His Ala Asp Thr Cys His Phe Asp
 35 40 45
 Val Asn Val Trp Glu Ala Ser Gly Asn Arg Ser Gly Gly Val Cys Asp
 50 55 60
 Asp Cys Gln His Asn Thr Glu Gly Gln Tyr Cys Gln Arg Cys Lys Pro
 65 70 75 80
 Gly Phe Tyr Arg Asp Leu Arg Arg Pro Phe Ser Ala Pro Asp Ala Cys
 85 90 95
 Lys Pro Cys Ser Cys His Pro Val Gly Ser Ala Val Leu Pro Ala Asn
 100 105 110

Ser Val Thr Phe Cys Asp Pro Ser Asn Gly Asp Cys Pro Cys Lys Pro
 115 120 125
 Gly Val Ala Gly Arg Arg Cys Asp Arg Cys Met Val Gly Tyr Trp Gly
 130 135 140
 Phe Gly Asp Tyr Gly Cys Arg Pro Cys Asp Cys Ala Gly Ser Cys Asp
 145 150 155 160
 Pro Ile Thr Gly Asp Cys Ile Ser Ser His Thr Asp Ile Asp Trp Tyr
 165 170 175
 His Glu Val Pro Asp Phe Arg Pro Val His Asn Lys Ser Glu Pro Ala
 180 185 190
 Trp Glu Trp Glu Asp Ala Gln Gly Phe Ser Ala Leu Leu His Ser Gly
 195 200 205
 Lys Cys Glu Cys Lys Glu Gln Thr Leu Gly Asn Ala Lys Ala Phe Cys
 210 215 220
 Gly Met Lys Tyr Ser Tyr Val Leu Lys Ile Lys Ile Leu Ser Ala His
 225 230 235 240
 Asp Lys Gly Thr His Val Glu Val Asn Val Lys Ile Lys Lys Val Leu
 245 250 255
 Lys Ser Thr Lys Leu Lys Ile Phe Arg Gly Lys Ala Asn Ile Ile Ser
 260 265 270
 Arg Ile Met Asp Gly Gln Arg Met His Leu Ser Asn Pro Gln Ser Trp
 275 280 285
 Phe Gly Ile Pro Cys Ser Arg Thr
 290 295

<210> 287
 <211> 37
 <212> PRT
 <213> Homo sapiens

<400> 287
 Cys Asp Asp Cys Gln His Asn Thr Glu Gly Gln Tyr Cys Gln Arg Cys
 1 5 10 15
 Lys Pro Gly Phe Tyr Arg Asp Leu Arg Arg Pro Phe Ser Ala Pro Asp
 20 25 30
 Ala Cys Lys Pro Cys
 35

<210> 288
 <211> 36
 <212> PRT
 <213> Homo sapiens

<400> 288

Cys Pro Cys Lys Pro Gly Val Ala Gly Arg Arg Cys Asp Arg Cys Met
 1 5 10 15

Val Gly Tyr Trp Gly Phe Gly Asp Tyr Gly Cys Arg Pro Cys Asp Cys
 20 25 30

Ala Gly Ser Cys
 35

<210> 289

<211> 66

<212> PRT

<213> Homo sapiens

<400> 289

Asn Ile Ser Ser Gln Tyr Cys Ile Leu Lys Ser Leu Glu Met Met Ile
 1 5 10 15

Ser Gly Leu Lys Leu Leu Val Leu Phe Leu Lys Phe Ala Pro Glu Asn
 20 25 30

Tyr Cys Leu Ser Thr Glu Thr Leu Gln Met Pro Asn Arg His Leu Arg
 35 40 45

Leu Ser Lys Ala Thr Cys Tyr Leu Met Lys Cys Leu Leu Pro Ser Tyr
 50 55 60

Phe Glu
 65

<210> 290

<211> 88

<212> PRT

<213> Homo sapiens

<400> 290

Pro Ile Glu Gly Thr Pro Ala Gly Thr Gly Pro Glu Phe Pro Gly Arg
 1 5 10 15

Pro Thr Arg Pro Gln Arg Met Arg Ser Leu Ile Ser Ser His Pro Cys
 20 25 30

Gln His Leu Leu Leu Leu Leu Leu Leu Phe Leu Ile Leu Ala Ile
 35 40 45

Leu Val Asp Val Lys Trp Tyr Leu Val Leu Phe Ile Cys Ile Ser Leu
 50 55 60

Met Thr Ser Asp Val Glu His Leu Phe Met Cys Leu Leu Ala Ile Arg
 65 70 75 80

Ile Ser Ser Trp Arg Asn Val Tyr
 85

<210> 291

<211> 60
<212> PRT
<213> Homo sapiens

<400> 291

Asn Trp Val Pro Thr Cys Leu Cys Pro Ser Ala Pro Cys Ser Phe His
1 5 10 15
Leu Leu Ser Arg Phe Lys Cys Leu Phe Ser Pro Gln Arg Leu Thr Asp
20 25 30
Ile Phe Arg Arg Tyr Asp Thr Asp Gln Asp Gly Trp Ile Gln Val Ser
35 40 45
Tyr Glu Gln Tyr Leu Ser Met Val Phe Ser Ile Val
50 55 60

<210> 292
<211> 33
<212> PRT
<213> Homo sapiens

<400> 292

Gln Arg Leu Thr Asp Ile Phe Arg Arg Tyr Asp Thr Asp Gln Asp Gly
1 5 10 15
Trp Ile Gln Val Ser Tyr Glu Gln Tyr Leu Ser Met Val Phe Ser Ile
20 25 30
Val

<210> 293
<211> 73
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (38)
<223> Xaa equals any of the naturally occurring L-amino acids

<220>
<221> SITE
<222> (48)
<223> Xaa equals any of the naturally occurring L-amino acids

<220>
<221> SITE
<222> (54)
<223> Xaa equals any of the naturally occurring L-amino acids

<220>
<221> SITE
<222> (55)
<223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (68)
 <223> Xaa equals any of the naturally occurring L-amino acids

 <400> 293
 Met Phe Tyr Lys Leu Thr Leu Ile Leu Cys Glu Leu Ser Val Ala Gly
 1 5 10 15
 Val Thr Gln Ala Ala Ser Gln Arg Pro Leu Gln Arg Leu Pro Arg His
 20 25 30
 Ile Cys Ser Gln Arg Xaa Pro Pro Gly Arg Cys Leu Leu Lys Ala Xaa
 35 40 45
 Leu Gln Thr Thr Trp Xaa Xaa Pro Asp Lys Pro Ile Pro Arg Leu Ser
 50 55 60
 Pro Pro Leu Xaa Ser Asp Pro Lys Arg
 65 70

<210> 294
 <211> 95
 <212> PRT
 <213> Homo sapiens

<400> 294
 Thr Ser Ser Pro Val Phe Ser Phe Cys Ser Met Ala Val Arg Glu Pro
 1 5 10 15
 Asp His Leu Gln Arg Val Ser Leu Pro Arg Tyr Asn Val Ser Ala Ser
 20 25 30
 Leu Gln Trp Leu Pro Cys His Arg Ile Val Leu Gln Pro Trp His Met
 35 40 45
 Cys Ala Met Trp Glu Leu Gly Gln Val Leu Phe His Pro Val Ala Pro
 50 55 60
 Arg Glu Gly Ala Ala Pro Ser Pro Val Ser Thr Leu Thr Trp Pro Ser
 65 70 75 80
 Ser Cys Ser His Ser Glu Ser Thr Met Glu Leu Glu Leu Gln Phe
 85 90 95

<210> 295
 <211> 16
 <212> PRT
 <213> Homo sapiens

<400> 295
 Met Ala Val Arg Glu Pro Asp His Leu Gln Arg Val Ser Leu Pro Arg
 1 5 10 15

<210> 296
 <211> 7
 <212> PRT
 <213> Homo sapiens

<400> 296
 Leu Pro Cys His Arg Ile Val
 1 5

<210> 297
 <211> 15
 <212> PRT
 <213> Homo sapiens

<400> 297
 Ser Leu Gln Trp Leu Pro Cys His Arg Ile Val Leu Gln Pro Trp
 1 5 10 15

<210> 298
 <211> 454
 <212> PRT
 <213> Homo sapiens

<400> 298
 Cys Phe Lys Arg Lys Pro Lys Arg Glu His Cys Ser Cys Pro Ile Thr
 1 5 10 15
 Tyr Gln Ser Leu Gly Asp Ile Leu Asn Ala Ser Phe Phe Ser Lys Arg
 20 25 30
 Lys Gly Met Gln Glu Val Lys Leu Asn Ser Tyr Val Val Ser Gly Thr
 35 40 45
 Ile Gly Leu Lys Glu Lys Ile Ser Leu Ser Glu Pro Val Phe Leu Thr
 50 55 60
 Phe Arg His Asn Gln Pro Gly Asp Lys Arg Thr Lys His Ile Cys Val
 65 70 75 80
 Tyr Trp Glu Gly Ser Glu Gly Gly Arg Trp Ser Thr Glu Gly Cys Ser
 85 90 95
 His Val His Ser Asn Gly Ser Tyr Thr Lys Cys Lys Cys Phe His Leu
 100 105 110
 Ser Ser Phe Ala Val Leu Val Ala Leu Ala Pro Lys Glu Asp Pro Val
 115 120 125
 Leu Thr Val Ile Thr Gln Val Gly Leu Thr Ile Ser Leu Leu Cys Leu
 130 135 140
 Phe Leu Ala Ile Leu Thr Phe Leu Leu Cys Arg Pro Ile Gln Asn Thr
 145 150 155 160
 Ser Thr Ser Leu His Leu Glu Leu Ser Leu Cys Leu Phe Leu Ala His
 165 170 175

Leu Leu Phe Leu Thr Gly Ile Asn Arg Thr Glu Pro Glu Val Leu Cys
 180 185 190
 Ser Ile Ile Ala Gly Leu Leu His Phe Leu Tyr Leu Ala Cys Phe Thr
 195 200 205
 Trp Met Leu Leu Glu Gly Leu His Leu Phe Leu Thr Val Arg Asn Leu
 210 215 220
 Lys Val Ala Asn Tyr Thr Ser Thr Gly Arg Phe Lys Lys Arg Phe Met
 225 230 235 240
 Tyr Pro Val Gly Tyr Gly Ile Pro Ala Val Ile Ile Ala Val Ser Ala
 245 250 255
 Ile Val Gly Pro Gln Asn Tyr Gly Thr Phe Thr His Cys Trp Leu Lys
 260 265 270
 Leu Asp Lys Gly Phe Ile Trp Ser Phe Met Gly Pro Val Ala Val Ile
 275 280 285
 Ile Leu Ile Asn Leu Val Phe Tyr Phe Gln Val Leu Trp Ile Leu Arg
 290 295 300
 Ser Lys Leu Ser Ser Leu Asn Lys Glu Val Ser Thr Ile Gln Asp Thr
 305 310 315 320
 Arg Val Met Thr Phe Lys Ala Ile Ser Gln Leu Phe Ile Leu Gly Cys
 325 330 335
 Ser Trp Gly Leu Gly Phe Phe Met Val Glu Glu Val Gly Lys Thr Ile
 340 345 350
 Gly Ser Ile Ile Ala Tyr Ser Phe Thr Ile Ile Asn Thr Leu Gln Gly
 355 360 365
 Val Leu Leu Phe Val Val His Cys Leu Leu Asn Arg Gln Val Arg Met
 370 375 380
 Glu Tyr Lys Lys Trp Phe Ser Gly Met Arg Lys Gly Val Glu Thr Glu
 385 390 395 400
 Ser Thr Glu Met Ser Arg Ser Thr Thr Gln Thr Lys Thr Glu Glu Val
 405 410 415
 Gly Lys Ser Ser Glu Ile Phe His Lys Gly Gly Thr Ala Ser Ser Ser
 420 425 430
 Ala Glu Ser Thr Lys Gln Pro Gln Pro Gln Val His Leu Val Ser Ala
 435 440 445
 Ala Trp Leu Lys Met Asn
 450

<210> 299

<211> 101

<212> PRT

<213> Homo sapiens

<400> 299

Phe Phe Trp Lys Glu Asn Leu Arg Arg Asn Gly Ser Arg Glu Asp Phe
1 5 10 15
Ala Arg Arg Ala Thr Gln Leu Ile Gln Ser Val Glu Leu Ser Ile Trp
20 25 30
Asn Ala Ser Phe Ala Ser Pro Gly Lys Gly Gln Ile Ser Glu Phe Asp
35 40 45
Ile Val Tyr Glu Thr Lys Arg Cys Asn Glu Thr Arg Glu Asn Ala Phe
50 55 60
Leu Glu Ala Gly Asn Asn Thr Met Asp Ile Asn Cys Ala Asp Ala Leu
65 70 75 80
Lys Gly Asn Leu Arg Glu Ser Thr Ala Val Ala Leu Ser Leu Ile Asn
85 90 95
Leu Leu Gly Ile Phe
100

<210> 300

<211> 27

<212> PRT

<213> Homo sapiens

<400> 300

Asp Ile Asn Glu Cys Glu Thr Gly Leu Ala Lys Cys Lys Tyr Lys Ala
1 5 10 15
Tyr Cys Arg Asn Lys Val Gly Gly Tyr Ile Cys
20 25

<210> 301

<211> 12

<212> PRT

<213> Homo sapiens

<400> 301

Cys Arg Asn Lys Val Gly Gly Tyr Ile Cys Ser Cys
1 5 10

<210> 302

<211> 331

<212> PRT

<213> Homo sapiens

<400> 302

Ala Leu Cys Pro His Pro His Leu Ile Leu Asn Val Thr Val Ser Pro
1 5 10 15
Ala Pro Ser Cys Arg His Val Lys Lys Val Val Ala Ser Pro Ser Pro
20 25 30

Ser Thr Thr Met Ile Ala Met Asp Ala Pro His Ser Lys Ala Ala Leu
 35 40 45
 Asp Ser Ile Asn Glu Leu Pro Glu Asn Ile Leu Leu Glu Leu Phe Thr
 50 55 60
 His Val Pro Ala Arg Gln Leu Leu Leu Asn Cys Arg Leu Val Cys Ser
 65 70 75 80
 Leu Trp Arg Asp Leu Ile Asp Leu Met Thr Leu Trp Lys Arg Lys Cys
 85 90 95
 Leu Arg Glu Gly Phe Ile Thr Lys Asp Trp Asp Gln Pro Val Ala Asp
 100 105 110
 Trp Lys Ile Phe Tyr Phe Leu Arg Ser Leu His Arg Asn Leu Leu Arg
 115 120 125
 Asn Pro Cys Ala Glu Glu Asp Met Phe Ala Trp Gln Ile Asp Phe Asn
 130 135 140
 Gly Gly Asp Arg Trp Lys Val Glu Ser Leu Pro Gly Ala His Gly Thr
 145 150 155 160
 Asp Phe Pro Asp Pro Lys Val Lys Lys Tyr Phe Val Thr Ser Tyr Glu
 165 170 175
 Met Cys Leu Lys Ser Gln Leu Val Asp Leu Val Ala Glu Gly Tyr Trp
 180 185 190
 Glu Glu Leu Leu Asp Thr Phe Arg Pro Asp Ile Val Val Lys Asp Trp
 195 200 205
 Phe Ala Ala Arg Ala Asp Cys Gly Cys Thr Tyr Gln Leu Lys Val Gln
 210 215 220
 Leu Ala Ser Ala Asp Tyr Phe Val Leu Ala Ser Phe Glu Pro Pro Pro
 225 230 235 240
 Val Thr Ile Gln Gln Trp Asn Asn Ala Thr Trp Thr Glu Val Ser Tyr
 245 250 255
 Thr Phe Ser Asp Tyr Pro Arg Gly Val Arg Tyr Ile Leu Phe Gln His
 260 265 270
 Gly Gly Arg Asp Thr Gln Tyr Trp Ala Gly Trp Tyr Gly Pro Arg Val
 275 280 285
 Thr Asn Ser Ser Ile Val Val Ser Pro Lys Met Thr Arg Asn Gln Ala
 290 295 300
 Ser Ser Glu Ala Gln Pro Gly Gln Lys His Gly Gln Glu Glu Ala Ala
 305 310 315 320
 Gln Ser Pro Tyr Arg Ala Val Val Gln Ile Phe
 325 330

<210> 303
 <211> 328
 <212> PRT
 <213> Homo sapiens

<400> 303

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Arg Gln Arg Ser Trp Asn Pro Gly Thr Asn Cys Tyr His Pro Asn Met
  1          5          10          15

Pro Asp Ala Phe Leu Thr Cys Glu Thr Val Ile Phe Ala Trp Ala Ile
      20          25          30

Gly Gly Glu Gly Phe Ser Tyr Pro Pro His Val Gly Leu Ser Leu Gly
  35          40          45

Thr Pro Leu Asp Pro His Tyr Val Leu Leu Glu Val His Tyr Asp Asn
  50          55          60

Pro Thr Tyr Glu Glu Gly Leu Ile Asp Asn Ser Gly Leu Arg Leu Phe
  65          70          75          80

Tyr Thr Met Asp Ile Arg Lys Tyr Asp Ala Gly Val Ile Glu Ala Gly
      85          90          95

Leu Trp Val Ser Leu Phe His Thr Ile Pro Pro Gly Met Pro Glu Phe
  100          105          110

Gln Ser Glu Gly His Cys Thr Leu Glu Cys Leu Glu Glu Ala Leu Glu
  115          120          125

Ala Glu Lys Pro Ser Gly Ile His Val Phe Ala Val Leu Leu His Ala
  130          135          140

His Leu Ala Gly Arg Gly Ile Arg Leu Arg His Phe Arg Lys Gly Lys
  145          150          155          160

Glu Met Lys Leu Leu Ala Tyr Asp Asp Phe Asp Phe Asn Phe Gln
  165          170          175

Glu Phe Gln Tyr Leu Lys Glu Glu Gln Thr Ile Leu Pro Gly Asp Asn
  180          185          190

Leu Ile Thr Glu Cys Arg Tyr Asn Thr Lys Asp Arg Ala Glu Met Thr
  195          200          205

Trp Gly Gly Leu Ser Thr Arg Ser Glu Met Cys Leu Ser Tyr Leu Leu
  210          215          220

Tyr Tyr Pro Arg Ile Asn Leu Thr Arg Cys Ala Ser Ile Pro Asp Ile
  225          230          235          240

Met Glu Gln Leu Gln Phe Ile Gly Val Lys Glu Ile Tyr Arg Pro Val
  245          250          255

Thr Thr Trp Pro Phe Ile Ile Lys Ser Pro Lys Gln Tyr Lys Asn Leu
  260          255          270

Ser Phe Met Asp Ala Met Asn Lys Phe Lys Trp Thr Lys Lys Glu Gly
  275          280          285

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Leu Ser Phe Asn Lys Leu Val Leu Ser Leu Pro Val Asn Val Arg Cys
 290 295 300

Ser Lys Thr Asp Asn Ala Glu Trp Ser Ile Pro Arg Asn Asp Ser Ile
 305 310 315 320

Thr Ser Arg Tyr Arg Lys Thr Leu
 325

<210> 304

<211> 272

<212> PRT

<213> Homo sapiens

<400> 304

Met Cys Cys Trp Pro Leu Leu Leu Leu Trp Gly Leu Leu Pro Gly Thr
 1 5 10 15

Ala Ala Gly Gly Ser Gly Arg Thr Tyr Pro His Arg Thr Leu Leu Asp
 20 25 30

Ser Glu Gly Lys Tyr Trp Leu Gly Trp Ser Gln Arg Gly Ser Gln Ile
 35 40 45

Ala Phe Arg Leu Gln Val Arg Thr Ala Gly Tyr Val Gly Phe Gly Phe
 50 55 60

Ser Pro Thr Gly Ala Met Ala Ser Ala Asp Ile Val Val Gly Gly Val
 65 70 75 80

Ala His Gly Arg Pro Tyr Leu Gln Asp Tyr Phe Thr Asn Ala Asn Arg
 85 90 95

Glu Leu Lys Lys Asp Ala Gln Gln Asp Tyr His Leu Glu Tyr Ala Met
 100 105 110

Glu Asn Ser Thr His Thr Ile Ile Glu Phe Thr Arg Glu Leu His Thr
 115 120 125

Cys Asp Ile Asn Asp Lys Ser Ile Thr Asp Ser Thr Val Arg Val Ile
 130 135 140

Trp Ala Tyr His His Glu Asp Ala Gly Glu Ala Gly Pro Lys Tyr His
 145 150 155 160

Asp Ser Asn Arg Gly Thr Lys Ser Leu Arg Leu Leu Asn Pro Glu Lys
 165 170 175

Thr Ser Val Leu Ser Thr Ala Leu Pro Tyr Phe Asp Leu Val Asn Gln
 180 185 190

Asp Val Pro Ile Pro Asn Lys Asp Thr Thr Tyr Trp Cys Gln Met Phe
 195 200 205

Lys Ile Pro Val Phe Gln Glu Lys His His Val Ile Lys Val Glu Pro
 210 215 220

Val Ile Gln Arg Gly His Glu Ser Leu Val His His Ile Leu Leu Tyr
 225 230 235 240
 Gln Cys Ser Asn Asn Phe Asn Asp Ser Val Pro Gly Ile Arg Ala Arg
 245 250 255
 Ile Ala Ile Thr Pro Thr Cys Pro Met His Ser Ser Pro Val Lys Leu
 260 265 270

<210> 305
 <211> 207
 <212> PRT
 <213> Homo sapiens

<400> 305
 Thr Gly Thr Phe Trp Ser Pro Arg Ser Gln Arg Arg Gly Cys Cys Gly
 1 5 10 15
 Arg Arg Ala Pro Arg Pro Glu Ala Met Glu Asn Gly Ala Val Tyr Ser
 20 25 30
 Pro Thr Thr Glu Glu Asp Pro Gly Pro Ala Arg Gly Pro Arg Ser Gly
 35 40 45
 Leu Ala Ala Tyr Phe Phe Met Gly Arg Leu Pro Leu Leu Arg Arg Val
 50 55 60
 Leu Lys Gly Leu Gln Leu Leu Ser Leu Leu Ala Phe Ile Cys Glu
 65 70 75 80
 Glu Val Val Ser Gln Cys Thr Leu Cys Gly Gly Leu Tyr Phe Phe Glu
 85 90 95
 Phe Val Ser Cys Ser Ala Phe Leu Leu Ser Leu Leu Ile Leu Ile Val
 100 105 110
 Tyr Cys Thr Pro Phe Tyr Glu Arg Val Asp Thr Thr Lys Val Lys Ser
 115 120 125
 Ser Asp Phe Tyr Ile Thr Leu Gly Thr Gly Cys Val Phe Leu Leu Ala
 130 135 140
 Ser Ile Ile Phe Val Ser Thr His Asp Arg Thr Ser Ala Glu Ile Ala
 145 150 155 160
 Ala Ile Val Phe Gly Phe Ile Ala Ser Phe Met Phe Leu Leu Asp Phe
 165 170 175
 Ile Thr Met Leu Tyr Glu Lys Arg Gln Glu Ser Gln Leu Arg Lys Pro
 180 185 190
 Glu Asn Thr Thr Arg Ala Glu Ala Leu Thr Glu Pro Leu Asn Ala
 195 200 205

<210> 306
 <211> 135
 <212> PRT
 <213> Homo sapiens

<400> 306
 Ala Ser Ala Pro Arg Val Met Arg Gly His Leu Ala Gly Phe Pro Ala
 1 5 10 15
 Leu Ser Gly Leu Ala Ser Val Cys Leu Trp Ala Thr Phe Ser Ala Gln
 20 25 30
 Leu Pro Gly Pro Val Ala Ala Thr Ser Trp Thr Pro Ala Pro Leu Gly
 35 40 45
 Cys Ser Ala Ala Arg Ser Gly Pro Glu Lys Arg Leu Gly Thr Ala Ala
 50 55 60
 Pro Gly Ser Ala Ala Ser Leu Ala Gln Ala Gly Pro Gly Ala Pro Cys
 65 70 75 80
 Arg Val Leu Pro Val Asp Pro Ala Pro Ala Ala Leu Asn Val Arg Glu
 85 90 95
 Pro Gly Trp Leu Gly Gly Leu Phe Asp Gly Ala Leu Leu Gln Val Leu
 100 105 110
 Leu Asn Phe Leu Arg Lys Ser Thr Asp Val Leu Met Asp Thr Arg Glu
 115 120 125
 Ala Glu Ser Leu Glu Val Glu
 130 135

<210> 307
 <211> 188
 <212> PRT
 <213> Homo sapiens

<400> 307
 Asn Lys Leu His Ser Phe Pro Val Phe Leu Ser Gln Leu Leu Leu Asp
 1 5 10 15
 Arg Gln Leu Leu His Ala Pro Gln Thr Leu Pro Thr Pro His Cys Gly
 20 25 30
 Gly Ser Ser Arg Pro Gly Pro Ser His Pro Pro Trp Leu Leu Ile Gln
 35 40 45
 Leu Pro Cys Val His Val Ala Leu Trp Gln Met Leu Arg Asp Phe Ser
 50 55 60
 Asp Ser Arg Ile Thr Pro Ser Thr Leu Thr Thr Gln Pro Ala Ala Gln
 65 70 75 80
 Thr Ala Ala Pro Ala Lys Asp Gln Glu Ser Asp Ile Val Gly Gly Glu
 85 90 95
 Gly Ile Leu Cys Asp Ile Ala Phe Leu Gln Glu Asp His Pro Leu Gly

100 105 110
Val Gly Gly Ala Ser Ala Pro Ser Ser Arg Arg Glu Leu Ser Arg Arg
115 120 125
Gly Val His Thr Gln Thr Leu Pro Glu Asp Gly Thr Leu His Gly Thr
130 135 140
Pro Ser Ser Ser Phe Asp Cys Gly Ile Lys Tyr Ile Ile Ser Trp Pro
145 150 155 160
Leu Ala Pro Gly Cys Asp Leu Pro Ser Leu Glu Leu Ser Leu Val Cys
165 170 175
Lys Gly Val Ser Ser Cys Met Gly Phe Ala Ala Gly
180 185

<210> 308
<211> 78
<212> PRT
<213> Homo sapiens

<400> 308
Pro Gly Arg Pro Thr Arg Pro Thr Lys Asn Lys Val Cys Val Cys Leu
1 5 10 15
Gly Met Leu Phe Trp Ala Tyr Pro Ile Cys Val Phe Ile Asp Ser Leu
20 25 30
Ser Cys Gln Pro Cys Leu Trp Ser Thr Gly Ala Thr Ser His Phe Asn
35 40 45
Ser Pro Thr Thr Ser Pro Leu Phe Thr Leu Phe Met Pro Cys Ala Leu
50 55 60
Ala Pro Asn Pro Phe Thr Gln Leu Gly Lys Leu Asp Asp Arg
65 70 75

<210> 309
<211> 10
<212> PRT
<213> Homo sapiens

<400> 309
Pro Val Asp Leu Thr Lys Thr Arg Leu Gln
1 5 10

<210> 310
<211> 10
<212> PRT
<213> Homo sapiens

<400> 310
Pro Thr Asp Val Leu Lys Ile Arg Met Gln
1 5 10

<210> 311
 <211> 313
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (117)
 <223> Xaa equals any of the naturally occurring L-amino acids

<400> 311
 Met Thr Phe Gly Ser Thr Ile Ser Pro Thr Ser Thr His Ala Ser Pro
 1 5 10 15
 Ser Leu Gly Phe Cys Cys Ser Trp Leu Leu Glu Asp Leu Glu Glu Gln
 20 25 30
 Leu Tyr Cys Ser Ala Phe Glu Glu Ala Ala Leu Thr Arg Arg Ile Cys
 35 40 45
 Asn Pro Thr Ser Cys Trp Leu Pro Leu Asp Met Glu Leu Leu His Arg
 50 55 60
 Gln Val Leu Ala Leu Gln Thr Gln Arg Val Leu Leu Gly Met Trp Leu
 65 70 75 80
 Arg Arg Ala Trp Asp Thr Trp Val Ser Pro Arg Arg Val Ala Pro Gly
 85 90 95
 Ser Arg Cys Leu Leu Thr Ala Ser His Pro Cys Thr Glu Lys Arg Arg
 100 105 110
 Lys Ala Ser Ala Xaa Gln Arg Asn Leu Gly Tyr Pro Leu Ala Met Leu
 115 120 125
 Cys Leu Leu Val Leu Thr Gly Leu Ser Val Leu Ile Val Ala Ile His
 130 135 140
 Ile Leu Glu Leu Leu Ile Asp Glu Ala Ala Met Pro Arg Gly Met Gln
 145 150 155 160
 Gly Thr Ser Leu Gly Gln Val Ser Phe Ser Lys Leu Gly Ser Phe Gly
 165 170 175
 Ala Val Ile Gln Val Val Leu Ile Phe Tyr Leu Met Val Ser Ser Val
 180 185 190
 Val Gly Phe Tyr Ser Ser Pro Leu Phe Arg Ser Leu Arg Pro Arg Trp
 195 200 205
 His Asp Thr Ala Met Thr Gln Ile Ile Gly Asn Cys Val Cys Leu Leu
 210 215 220
 Val Leu Ser Ser Ala Leu Pro Val Phe Ser Arg Thr Leu Gly Leu Thr
 225 230 235 240
 Arg Phe Asp Leu Leu Gly Asp Phe Gly Arg Phe Asn Trp Leu Gly Asn
 245 250 255

Phe Tyr Ile Val Phe Leu Tyr Asn Ala Ala Phe Ala Gly Leu Thr Thr
 260 265 270
 Leu Cys Leu Val Lys Thr Phe Thr Ala Ala Val Arg Ala Glu Leu Ile
 275 280 285
 Arg Ala Phe Gly Leu Asp Arg Leu Pro Leu Pro Val Ser Gly Phe Pro
 290 295 300
 Gln Ala Ser Arg Lys Thr Gln His Gln
 305 310

<210> 312
 <211> 92
 <212> PRT
 <213> Homo sapiens

<400> 312
 Leu Cys Val Cys Leu Val Tyr Leu Cys Met Tyr Gly Val Cys Leu Cys
 1 5 10 15
 Val Ile Val Cys Val Ser Gly Val Ser Leu Cys Leu Tyr Val Trp Gly
 20 25 30
 Val Ser Val Cys Asp Cys Val Ser Val Phe Met Cys Val Cys Leu Cys
 35 40 45
 Val Ile Phe Cys Val Tyr Gly Lys Pro Arg Thr Glu His Tyr His Ser
 50 55 60
 Pro His Leu Ala Lys Gln Lys Ala Phe Arg Glu Met Cys Gly Arg His
 65 70 75 80
 Asp Val Ser Ala Ala Gly Ile Phe Gln Ser Tyr Val
 85 90

<210> 313
 <211> 207
 <212> PRT
 <213> Homo sapiens

<400> 313
 Gly His Met Pro Tyr Gly Trp Leu Thr Glu Ile Arg Ala Val Tyr Pro
 1 5 10 15
 Ala Phe Asp Lys Asn Asn Pro Ser Asn Lys Leu Val Ser Thr Ser Asn
 20 25 30
 Thr Val Thr Ala Ala His Ile Lys Lys Phe Thr Phe Val Cys Met Ala
 35 40 45
 Leu Ser Leu Thr Leu Cys Phe Val Met Phe Trp Thr Pro Asn Val Ser
 50 55 60
 Glu Lys Ile Leu Ile Asp Ile Ile Gly Val Asp Phe Ala Phe Ala Glu
 65 70 75 80

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<210> 314
<211> 114
<212> PRT
<213> Homo sapiens
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<210> 315
<211> 115

<212> PRT

<213> Homo sapiens

<400> 315

Arg Cys Cys Cys Arg Gly Cys Ser Cys Arg Ala Arg Leu Cys Pro Pro
1 5 10 15
Ala Arg Ser Thr Ala Val Ala Pro Glu Cys Arg Gly Ala His Pro Ser
20 25 30
Arg Ala Met Arg Pro Gly Thr Ala Leu Gln Ala Val Leu Leu Ala Val
35 40 45
Leu Leu Val Gly Leu Arg Ala Ala Thr Gly Arg Leu Leu Ser Gly Gln
50 55 60
Pro Val Cys Arg Gly Gly Thr Gln Arg Pro Cys Tyr Lys Val Ile Tyr
65 70 75 80
Phe His Asp Thr Ser Arg Arg Leu Asn Phe Glu Glu Ala Lys Glu Ala
85 90 95
Cys Arg Arg Gly Trp Arg Pro Ala Ser Gln His Arg Val Leu Lys Met
100 105 110
Asn Arg Asn
115

<210> 316

<211> 81

<212> PRT

<213> Homo sapiens

<400> 316

Met Arg Pro Gly Thr Ala Leu Gln Ala Val Leu Leu Ala Val Leu Leu
1 5 10 15
Val Gly Leu Arg Ala Ala Thr Gly Arg Leu Leu Ser Gly Gln Pro Val
20 25 30
Cys Arg Gly Gly Thr Gln Arg Pro Cys Tyr Lys Val Ile Tyr Phe His
35 40 45
Asp Thr Ser Arg Arg Leu Asn Phe Glu Glu Ala Lys Glu Ala Cys Arg
50 55 60
Arg Gly Trp Arg Pro Ala Ser Gln His Arg Val Leu Lys Met Asn Arg
65 70 75 80
Asn

<210> 317

<211> 290

<212> PRT

<213> Homo sapiens

<400> 317
 Ile Arg His Glu Gln Gln Gly Glu Glu Asp Asp Glu His Ala Arg Pro
 1 5 10 15
 Leu Ala Glu Ser Leu Leu Leu Ala Ile Ala Asp Leu Leu Phe Cys Pro
 20 25 30
 Asp Phe Thr Val Gln Ser His Arg Arg Ser Thr Val Asp Ser Ala Glu
 35 40 45
 Asp Val His Ser Leu Asp Ser Cys Glu Tyr Ile Trp Glu Ala Gly Val
 50 55 60
 Gly Phe Ala His Ser Pro Gln Pro Asn Tyr Ile His Asp Met Asn Arg
 65 70 75 80
 Met Glu Leu Leu Lys Leu Leu Leu Thr Cys Phe Ser Glu Ala Met Tyr
 85 90 95
 Leu Pro Pro Ala Pro Glu Ser Gly Ser Thr Asn Pro Trp Val Gln Phe
 100 105 110
 Phe Cys Ser Thr Glu Asn Arg His Ala Leu Pro Leu Phe Thr Ser Leu
 115 120 125
 Leu Asn Thr Val Cys Ala Tyr Asp Pro Val Gly Tyr Gly Ile Pro Tyr
 130 135 140
 Asn His Leu Leu Phe Ser Asp Tyr Arg Glu Pro Leu Val Glu Glu Ala
 145 150 155 160
 Ala Gln Val Leu Ile Val Thr Leu Asp His Asp Ser Ala Ser Ser Ala
 165 170 175
 Ser Pro Thr Val Asp Gly Thr Thr Thr Gly Thr Ala Met Asp Asp Ala
 180 185 190
 Asp Pro Pro Gly Pro Glu Asn Leu Phe Val Asn Tyr Leu Ser Arg Ile
 195 200 205
 His Arg Glu Glu Asp Phe Gln Phe Ile Leu Lys Gly Ile Ala Arg Leu
 210 215 220
 Leu Ser Asn Pro Leu Leu Gln Thr Tyr Leu Pro Asn Ser Thr Lys Lys
 225 230 235 240
 Asp Pro Val Pro Pro Gly Ala Ala Ser Ser Leu Leu Glu Ala Leu Arg
 245 250 255
 Leu Gln Gln Glu Ile Pro Leu Leu Arg Ala Glu Glu Gln Arg Arg Pro
 260 265 270
 Arg His Pro Cys Pro His Pro Leu Leu Pro Gln Arg Cys Pro Gly Arg
 275 280 285
 Ser Val
 290

<210> 318
 <211> 318
 <212> PRT
 <213> Homo sapiens

<400> 318

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Arg Leu Val Tyr Asn Lys Thr Ser Arg Ala Thr Gln Phe Pro Asp Gly
 1           5           10           15

Val Asp Val Arg Val Pro Gly Phe Gly Lys Thr Phe Ser Leu Glu Phe
 20           25           30

Leu Asp Pro Ser Lys Ser Ser Val Gly Ser Tyr Phe His Thr Met Val
 35           40           45

Glu Ser Leu Val Gly Trp Gly Tyr Thr Arg Gly Glu Asp Val Arg Gly
 50           55           60

Ala Pro Tyr Asp Trp Arg Arg Ala Pro Asn Glu Asn Gly Pro Tyr Phe
 65           70           75           80

Leu Ala Leu Arg Glu Met Ile Glu Glu Met Tyr Gln Leu Tyr Gly Gly
 85           90           95

Pro Val Val Leu Val Ala His Ser Met Gly Asn Met Tyr Thr Leu Tyr
100           105           110

Phe Leu Gln Arg Gln Pro Gln Ala Trp Lys Asp Lys Tyr Ile Arg Ala
115           120           125

Phe Val Ser Leu Gly Ala Pro Trp Gly Gly Val Ala Lys Thr Leu Arg
130           135           140

Val Leu Ala Ser Gly Asp Asn Asn Arg Ile Pro Val Ile Gly Pro Leu
145           150           155           160

Lys Ile Arg Glu Gln Gln Arg Ser Ala Val Ser Thr Ser Trp Leu Leu
165           170           175

Pro Tyr Asn Tyr Thr Trp Ser Pro Glu Lys Val Phe Val Gln Thr Pro
180           185           190

Thr Ile Asn Tyr Thr Leu Arg Asp Tyr Arg Lys Phe Phe Gln Asp Ile
195           200           205

Gly Phe Glu Asp Gly Trp Leu Met Arg Gln Asp Thr Glu Gly Leu Val
210           215           220

Glu Ala Thr Met Pro Pro Gly Val Gln Leu His Cys Leu Tyr Gly Thr
225           230           235           240

Gly Val Pro Thr Pro Asp Ser Phe Tyr Tyr Glu Ser Phe Pro Asp Arg
245           250           255

Asp Pro Lys Ile Cys Phe Gly Asp Gly Asp Gly Thr Val Asn Leu Lys
260           265           270

Ser Ala Leu Gln Cys Gln Ala Trp Gln Ser Arg Gln Glu His Gln Val
275           280           285

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Leu Leu Gln Glu Leu Pro Gly Ser Glu His Ile Glu Met Leu Ala Asn
 290 295 300

Ala Thr Thr Leu Ala Tyr Leu Lys Arg Val Leu Leu Gly Pro
 305 310 315

<210> 319

<211> 362

<212> PRT

<213> Homo sapiens

<400> 319

Met Asn Lys Glu Asp Lys Val Trp Asn Asp Cys Lys Gly Val Asn Lys
 1 5 10 15

Leu Thr Asn Leu Glu Glu Gln Tyr Ile Ile Leu Ile Phe Gln Asn Gly
 20 25 30

Leu Asp Pro Pro Ala Asn Met Val Phe Glu Ser Ile Ile Asn Glu Ile
 35 40 45

Gly Ile Lys Asn Asn Ile Ser Asn Phe Phe Ala Lys Ile Pro Phe Glu
 50 55 60

Glu Ala Asn Gly Arg Leu Val Ala Cys Thr Arg Thr Tyr Glu Glu Ser
 65 70 75 80

Ile Lys Gly Ser Cys Gly Gln Lys Glu Asn Lys Ile Lys Thr Val Ser
 85 90 95

Phe Glu Ser Lys Ile Gln Leu Arg Ser Lys Gln Glu Phe Gln Phe Phe
 100 105 110

Asp Glu Glu Glu Glu Thr Gly Glu Asn His Thr Ile Phe Ile Gly Pro
 115 120 125

Val Glu Lys Leu Ile Val Tyr Pro Pro Pro Ala Lys Gly Gly Ile
 130 135 140

Ser Val Thr Asn Glu Asp Leu His Cys Leu Asn Glu Gly Glu Phe Leu
 145 150 155 160

Asn Asp Val Ile Ile Asp Phe Tyr Leu Lys Tyr Leu Val Leu Glu Lys
 165 170 175

Leu Lys Lys Glu Asp Ala Asp Arg Ile His Ile Phe Ser Ser Phe Phe
 180 185 190

Tyr Lys Arg Leu Asn Gln Arg Glu Arg Arg Asn His Glu Thr Thr Asn
 195 200 205

Leu Ser Ile Gln Gln Lys Arg His Gly Arg Val Lys Thr Trp Thr Arg
 210 215 220

His Val Asp Ile Phe Glu Lys Asp Phe Ile Phe Val Pro Leu Asn Glu
 225 230 235 240

Ala Ala His Trp Phe Leu Ala Val Val Cys Phe Pro Gly Leu Glu Lys
 245 250 255
 Pro Lys Tyr Glu Pro Asn Pro His Tyr His Glu Asn Ala Val Ile Gln
 260 265 270
 Lys Cys Ser Thr Val Glu Asp Ser Cys Ile Ser Ser Ser Ala Ser Glu
 275 280 285
 Met Glu Ser Cys Ser Gln Asn Ser Ser Ala Lys Pro Val Ile Lys Lys
 290 295 300
 Met Leu Asn Lys Lys His Cys Ile Ala Val Ile Asp Ser Asn Pro Gly
 305 310 315 320
 Gln Glu Glu Ser Asp Pro Arg Tyr Lys Arg Asn Ile Cys Ser Val Lys
 325 330 335
 Tyr Ser Val Lys Lys Ile Asn His Thr Ala Ser Glu Asn Glu Glu Phe
 340 345 350
 Asn Lys Gly Glu Ser Thr Ser Gln Lys Ser
 355 360

<210> 320

<211> 330

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (38)

<223> Xaa equals any of the naturally occurring L-amino acids

<220>

<221> SITE

<222> (247)

<223> Xaa equals any of the naturally occurring L-amino acids

<400> 320

Met Ser Pro Leu Ser Ala Ala Arg Ala Ala Leu Arg Val Tyr Ala Val
 1 5 10 15
 Gly Ala Ala Val Ile Leu Ala Gln Leu Leu Arg Arg Cys Arg Gly Gly
 20 25 30
 Phe Leu Glu Pro Val Xaa Pro Pro Arg Pro Asp Arg Val Ala Ile Val
 35 40 45
 Thr Gly Gly Thr Asp Gly Ile Gly Tyr Ser Thr Ala Asn Ile Trp Arg
 50 55 60
 Asp Leu Gly Met His Val Ile Ile Ala Gly Asn Asn Asp Ser Lys Ala
 65 70 75 80
 Lys Gln Val Val Ser Lys Ile Lys Glu Glu Thr Leu Asn Asp Lys Val
 85 90 95

Glu Phe Leu Tyr Cys Asp Leu Ala Ser Met Thr Ser Ile Arg Gln Phe
 100 105 110
 Val Gln Lys Phe Lys Met Lys Lys Ile Pro Leu His Val Leu Ile Asn
 115 120 125
 Asn Ala Gly Val Met Met Val Pro Gln Arg Lys Thr Arg Asp Gly Phe
 130 135 140
 Glu Glu His Phe Gly Leu Asn Tyr Leu Gly His Phe Leu Leu Thr Asn
 145 150 155 160
 Leu Leu Leu Asp Thr Leu Lys Glu Ser Gly Ser Pro Gly His Ser Ala
 165 170 175
 Arg Val Val Thr Val Ser Ser Ala Thr His Tyr Val Ala Glu Leu Asn
 180 185 190
 Met Asp Asp Leu Gln Ser Ser Ala Cys Tyr Ser Pro His Ala Ala Tyr
 195 200 205
 Ala Gln Ser Lys Leu Ala Leu Val Leu Phe Thr Tyr His Leu Gln Arg
 210 215 220
 Leu Leu Ala Ala Glu Gly Ser His Val Thr Ala Asn Val Val Asp Pro
 225 230 235 240
 Gly Val Val Asn Thr Asp Xaa Tyr Lys His Val Phe Trp Ala Thr Arg
 245 250 255
 Leu Ala Lys Lys Leu Leu Gly Trp Leu Leu Phe Lys Thr Pro Asp Glu
 260 265 270
 Gly Ala Trp Thr Ser Ile Tyr Ala Ala Val Thr Pro Glu Leu Glu Gly
 275 280 285
 Val Gly Gly Arg Tyr Leu Tyr Asn Glu Lys Glu Thr Lys Ser Leu His
 290 295 300
 Val Thr Tyr Asn Gln Lys Leu Gln Gln Gln Leu Trp Ser Lys Ser Cys
 305 310 315 320
 Glu Met Thr Gly Val Leu Asp Val Thr Leu
 325 330

<210> 321

<211> 71

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (38)

<223> Xaa equals any of the naturally occurring L-amino acids

<400> 321

Met Ser Pro Leu Ser Ala Ala Arg Ala Ala Leu Arg Val Tyr Ala Val
 1 5 10 15

Gly Ala Ala Val Ile Leu Ala Gln Leu Leu Arg Arg Cys Arg Gly Gly
 20 25 30
 Phe Leu Glu Pro Val Xaa Pro Pro Arg Pro Asp Arg Val Ala Ile Val
 35 40 45
 Thr Gly Gly Thr Asp Gly Ile Gly Tyr Ser Thr Ala Asn Ile Trp Arg
 50 55 60
 Asp Leu Ala Cys Met Leu Ser
 65 70

<210> 122
 <211> 266
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (97)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (174)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (195)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (199)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (206)
 <223> Xaa equals any of the naturally occurring L-amino acids

<400> 322
 Met Glu Val Thr Thr Glu Asp Thr Ser Arg Thr Asp Val Ser Glu Pro
 1 5 10 15
 Ala Thr Ser Gly Gly Ala Ala Asp Gly Val Thr Ser Ile Ala Pro Thr
 20 25 30
 Ala Val Ala Ser Ser Thr Thr Ala Ala Ser Ile Thr Thr Ala Ala Ser
 35 40 45
 Ser Met Thr Val Ala Ser Ser Ala Pro Thr Thr Ala Ala Ser Ser Thr
 50 55 60
 Thr Val Ala Ser Ile Ala Pro Thr Thr Thr Ala Ser Ser Met Thr Ala
 65 70 75 80

Ala Ser Ser Thr Pro Met Thr Leu Ala Leu Pro Ala Pro Thr Ser Thr
85 90 95

Xaa Thr Gly Arg Thr Pro Ser Thr Thr Ala Thr Gly His Pro Ser Leu
100 105 110

Ser Thr Ala Leu Ala Gln Val Pro Lys Ser Ser Ala Leu Pro Arg Thr
115 120 125

Ala Thr Leu Ala Thr Leu Ala Thr Arg Ala Gln Thr Val Ala Thr Thr
130 135 140

Ala Asn Thr Ser Ser Pro Met Ser Thr Arg Pro Ser Pro Ser Lys His
145 150 155 160

Met Pro Ser Asp Thr Ala Ala Ser Pro Val Pro Pro Met Xaa Pro Gln
165 170 175

Ala Gln Gly Pro Ile Ser Gln Val Ser Val Asp Gln Pro Val Val Asn
180 185 190

Thr Thr Xaa Lys Ser Thr Xaa Met Pro Ser Asn Thr Thr Xaa Glu Pro
195 200 205

Leu Thr Gln Ala Val Val Asp Lys Thr Leu Leu Leu Val Val Leu Leu
210 215 220

Leu Gly Val Thr Leu Phe Ile Thr Val Leu Val Leu Phe Ala Leu Gln
225 230 235 240

Ala Tyr Glu Ser Tyr Lys Lys Lys Asp Tyr Thr Gln Val Asp Tyr Leu
245 250 255

Ile Asn Gly Met Tyr Ala Asp Ser Glu Met
260 265

<210> 323

<211> 99

<212> PRT

<213> Homo sapiens

<400> 323

Ala Arg Cys Pro Glu Leu Pro Gly Leu Arg Cys Arg Pro Arg Pro Arg
1 5 10 15

Ala Gly Pro Gln Ala Pro Ser Tyr Cys Pro Arg Ala Thr Arg Pro Pro
20 25 30

Gly Ala Cys Cys Ala Arg Met Arg Leu Leu Leu Glu Trp Arg Val Tyr
35 40 45

Leu Arg Leu Thr Cys Ala Thr Lys Asp Gly Met Ala Arg Glu Cys Pro
50 55 60

Thr Thr Trp Leu Ser Pro Ala Lys Pro Asp Phe Ala Gln Arg His
65 70 75 80

Ser Val Lys Pro Thr Ala Leu Gln Gly Gly Arg Trp Ser Arg Leu Gly
 85 90 95

Ala Ser Pro

<210> 324

<211> 96

<212> PRT

<213> Homo sapiens

<400> 324

Leu Pro Ala Thr Val Glu Phe Ala Val His Thr Phe Asn Gln Gln Ser
 1 5 10 15

Lys Asp Tyr Tyr Ala Tyr Arg Leu Gly His Ile Leu Asn Ser Trp Lys
 20 25 30

Glu Gln Val Glu Ser Lys Thr Val Phe Ser Met Glu Leu Leu Leu Gly
 35 40 45

Arg Thr Arg Cys Gly Lys Phe Glu Asp Asp Ile Asp Asn Cys His Phe
 50 55 60

Gln Glu Ser Thr Glu Leu Asn Asn Thr Phe Thr Cys Phe Phe Thr Ile
 65 70 75 80

Ser Thr Arg Pro Trp Met Thr Gln Phe Ser Leu Leu Asn Lys Thr Cys
 85 90 95

<210> 325

<211> 166

<212> PRT

<213> Homo sapiens

<400> 325

Leu Leu Trp Ala Arg Gly Leu Gly Arg Ala Lys Ser Ala Val Pro Thr
 1 5 10 15

Val Ser Thr Met Leu Gly Leu Pro Trp Lys Gly Gly Leu Ser Trp Ala
 20 25 30

Leu Leu Leu Leu Leu Leu Gly Ser Gln Ile Leu Leu Ile Tyr Ala Trp
 35 40 45

His Phe His Glu Gln Arg Asp Cys Asp Glu His Asn Val Met Ala Arg
 50 55 60

Tyr Leu Pro Ala Thr Val Glu Phe Ala Val His Thr Phe Asn Gln Gln
 65 70 75 80

Ser Lys Asp Tyr Tyr Ala Tyr Arg Leu Gly His Ile Leu Asn Ser Trp
 85 90 95

Lys Glu Gln Val Glu Ser Lys Thr Val Phe Ser Met Glu Leu Leu Leu
 100 105 110
 Gly Arg Thr Arg Cys Gly Lys Phe Glu Asp Asp Ile Asp Asn Cys His
 115 120 125
 Phe Gln Glu Ser Thr Glu Leu Asn Asn Thr Phe Thr Cys Phe Phe Thr
 130 135 140
 Ile Ser Thr Arg Pro Trp Met Thr Gln Phe Ser Leu Leu Asn Lys Thr
 145 150 155 160
 Cys Leu Glu Gly Phe His
 165

<210> 326

<211> 214

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (200)

<223> Xaa equals any of the naturally occurring L-amino acids

<220>

<221> SITE

<222> (205)

<223> Xaa equals any of the naturally occurring L-amino acids

<400> 326

Leu Glu Gln Lys Leu Glu Leu His Arg Gly Gly Gly Arg Ser Arg Thr
 1 5 10 15
 Ser Gly Ser Pro Gly Leu Gln Glu Phe Gly Thr Arg Glu Glu Arg Gly
 20 25 30
 Glu Gly Glu Gln Arg Thr Gly Arg Glu Phe Ser Gly Asn Gly Gly Arg
 35 40 45
 Ala Val Glu Ala Ala Arg Met Arg Leu Leu Cys Gly Leu Trp Leu Trp
 50 55 60
 Leu Ser Leu Leu Lys Val Leu Gln Ala Gln Thr Pro Thr Pro Leu Pro
 65 70 75 80
 Leu Pro Pro Pro Met Gln Ser Phe Gln Gly Asn Gln Phe Gln Gly Glu
 85 90 95
 Trp Phe Val Leu Gly Leu Ala Gly Asn Ser Phe Arg Pro Glu His Arg
 100 105 110
 Ala Leu Leu Asn Ala Phe Thr Ala Thr Phe Glu Leu Ser Asp Asp Gly
 115 120 125
 Arg Phe Glu Val Trp Asn Ala Met Thr Arg Gly Gln His Cys Asp Thr
 130 135 140

Trp Ser Tyr Val Leu Ile Pro Ala Ala Gln Pro Gly Gln Phe Thr Val
 145 150 155 160
 Asp His Gly Val Gly Arg Ser Trp Leu Leu Pro Pro Gly Thr Leu Asp
 165 170 175
 Gln Phe Ile Cys Leu Gly Arg Ala Gln Gly Leu Ser Asp Asp Asn Ile
 180 185 190
 Val Phe Pro Asp Val Thr Gly Xaa Ala Leu Asp Leu Xaa Ser Leu Pro
 195 200 205
 Trp Val Ala Ala Pro Ala
 210

<210> 327
 <211> 181
 <212> PRT
 <213> Homo sapiens

<400> 327
 Met Cys Val Cys Glu Arg Lys Arg Gly Arg Glu Lys Glu Gly Gly Val
 1 5 10 15
 Thr Pro Thr Met Thr Ser Asn Phe Pro Phe Cys Thr Leu Ile Leu Gly
 20 25 30
 Ile Ala Gln Ala Gln Ala Cys Pro Gly Cys Pro Gly Asp Trp Pro Gly
 35 40 45
 Leu Gly Ser Gly Val Gly Glu Gly Leu His His Ile Arg Thr Cys Arg
 50 55 60
 Thr Pro Ile Pro Cys Ser Pro Pro Ala Pro Ala Ala Ala Cys Leu Gly
 65 70 75 80
 Ser Gly His Ala Arg Leu Pro Cys Val Leu Arg Leu Trp Pro Val Pro
 85 90 95
 Ala Asn Leu Ser Ser Pro Phe Arg Leu Glu Ala Leu His Cys Ser Phe
 100 105 110
 Trp Ser Ser Pro Leu Leu Pro Ala Pro His Leu Ala Phe Phe Gly Phe
 115 120 125
 Arg Asp Leu Leu Thr Asp Phe Leu Leu Ala Ala Cys Leu Leu Thr Phe
 130 135 140
 Gln Lys Thr Pro Leu Glu Leu Pro Met Ala Val Val His Leu Leu Val
 145 150 155 160
 Ala Thr Pro Cys Tyr Gln Met Leu Asp Asn Leu Pro Leu Pro Ser Ala
 165 170 175
 Ala Ala Asn Trp Cys
 180

<210> 328
 <211> 195
 <212> PRT
 <213> Homo sapiens

<400> 328
 Tyr Leu Trp Gly Arg Pro Arg Leu Arg Met Arg Ala Gly Thr Ser Pro
 1 5 10 15
 Ser Ala Pro Trp Gly Glu Lys Arg Glu Lys Leu Gly His Lys Leu Pro
 20 25 30
 Val Ala Leu Gln Gly Tyr His Pro Trp Ile Leu Leu Glu Cys Thr Val
 35 40 45
 Phe Trp Ala Arg Val Val Leu Ala Cys Phe Ser Leu Tyr Leu Ile Arg
 50 55 60
 Gly Pro Asn Cys Ile Asn Arg Gln Pro Glu Pro Thr Tyr Gln Lys Ala
 65 70 75 80
 Cys Asn Leu Asp Cys Ser Ser Asp Phe Gly Gln Glu Arg Ala Pro Ala
 85 90 95
 Trp Glu Leu Leu Gly Pro Glu Ser Glu Gln Arg Leu Arg Glu Tyr Thr
 100 105 110
 Ala Gln Gly Leu Gln Ser Leu Ala Ser Ser His Arg Trp Arg Gln Phe
 115 120 125
 Lys Thr Glu Gly Lys Met Arg Gly Gly Ala Ser Pro Leu Pro Trp Leu
 130 135 140
 Ile Cys Phe Trp Leu Cys Ser Tyr Lys Gly Ser Asp Asn Ser Leu Lys
 145 150 155 160
 Pro Val Val Pro Gly Pro Thr Leu Cys Pro Gln Ser Leu Val Ser Pro
 165 170 175
 Ser Val His Pro Ser Thr Arg Ser Ala Ser Leu Gly Arg His Arg Ala
 180 185 190
 Glu Ala Ala
 195

<210> 329
 <211> 50
 <212> PRT
 <213> Homo sapiens

<400> 329
 Met Pro Gly Ile Leu Ala Gly Ile Pro Val Lys Asp Leu Cys Leu Ser
 1 5 10 15
 Leu Leu Gln Gly Phe Arg Leu Leu Leu Cys Val Cys Pro Gly Trp
 20 25 30
 Leu Ser Gly Trp Met Gly Gly Gln Lys Gly Ser Pro Arg Ile Val Asp

35 40 45

Ile Gly
50

<210> 330
<211> 90
<212> PRT
<213> Homo sapiens

<400> 330
Ala Lys Gly Glu Glu Arg Lys Glu Ala Phe Ser Leu Lys Met Val Gln
1 5 10 15
Leu Ser Ser Glu Pro Ile Ser Phe Gly Leu Met Tyr Leu Tyr Leu Gly
20 25 30
Val Phe Phe His Leu Ile Tyr Pro Gly Ala Leu Ser Ile Thr Thr Leu
35 40 45
Gly Lys His Ser His Pro Phe Phe Thr Ala Glu Gln Asn Ser Thr Val
50 55 60
Trp Met Glu His Thr Leu Phe His Gln Ser Pro Val Ala Ser His Leu
65 70 75 80
Val Cys Phe Gln Ser Phe Ala Phe Ser Glu
85 90

<210> 331
<211> 56
<212> PRT
<213> Homo sapiens

<400> 331
Gly Pro Ala His Pro Ala Ser Pro Pro Leu Met Thr Leu Ser Leu Gln
1 5 10 15
Leu Ala Glu Leu Val His Phe Val Cys Ala Phe Gln Ser Gln Trp Thr
20 25 30
Gly Val Tyr Pro Met Met Pro Pro Leu Lys Pro Thr Glu Pro Leu Cys
35 40 45
Phe Ala Cys Val Pro Cys Arg Val
50 55

<210> 332
<211> 18
<212> PRT
<213> Homo sapiens

<400> 332
Met Leu Leu Glu Val Tyr Gly Asp Ser Ile Ser Val Thr Val Ala Ile
1 5 10 15

Pro Leu

<210> 333
 <211> 19
 <212> PRT
 <213> Homo sapiens

<400> 333
 Met His Ser Pro Cys Gln Ser Lys Ala Ala Asp Gly Leu Gly Lys Ser
 1 5 10 15

Glu Thr Glu

<210> 334
 <211> 10
 <212> PRT
 <213> Homo sapiens

<400> 334
 Met Leu Lys Ser Leu Gly Leu Ser Thr Asn
 1 5 10

<210> 335
 <211> 200
 <212> PRT
 <213> Homo sapiens

<400> 335
 Ala Gln Arg Leu Ala Glu Glu Cys Phe Tyr Met Leu Leu Glu Val Tyr
 1 5 10 15

Gly Asp Ser Ile Ser Val Thr Val Ala Ile Pro Leu Met His Ser Pro
 20 25 30

Cys Gln Ser Lys Ala Ala Asp Gly Leu Gly Lys Ser Glu Thr Glu Met
 35 40 45

Leu Lys Ser Leu Gly Leu Ser Thr Asn Met Ser Pro Phe His Leu Leu
 50 55 60

Gly Leu Lys Val Phe Leu Thr Trp Ala Leu Thr Leu Ala Gln Ile Cys
 65 70 75 80

Leu Tyr Phe Phe Glu Val Gln Pro Leu Gly Leu Leu Ala Leu Asn Phe
 85 90 95

Phe Cys Thr Ala Thr Ala Gly Leu Lys Glu Leu Cys Met His Pro Pro
 100 105 110

Ser Leu Ala Phe Thr Pro Glu Phe His Thr Ser Leu Ser Pro Leu Ala
 115 120 125

Ile Pro Ser Phe Cys Gly Thr Ser Val Ser Leu Ser Asn Ser His Thr
 130 135 140

Ile Pro Leu Ser Leu Tyr Leu Pro Phe Pro Ser Lys Ser Arg Met Pro
 145 150 155 160
 Asp Thr Leu His Leu Leu Val His Ser Leu Pro Leu Val His Ser Gln
 165 170 175
 Val Leu Pro Val Lys Asp Val Thr Ile Glu Trp Pro Leu Cys Gln Arg
 180 185 190
 Cys Leu Gly Ser Thr Cys His Gln
 195 200

<210> 336

<211> 99

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (94)

<223> Xaa equals any of the naturally occurring L-amino acids

<220>

<221> SITE

<222> (99)

<223> Xaa equals any of the naturally occurring L-amino acids

<400> 336

Trp Ile Pro Arg Ala Ala Gly Ile Arg His Glu Val Gln Val Ser Leu
 1 5 10 15
 Phe Gln Met Phe Cys Phe Ser Ser Ile Phe Cys Ser His Glu His Thr
 20 25 30
 His Leu Pro Gly Thr Phe Trp Leu Phe Leu Phe Leu Ile Leu
 35 40 45
 Pro Pro Ser Cys Pro Cys Phe Leu Pro Phe Ser Leu Ala Ile Glu Thr
 50 55 60
 Val Arg Trp Pro Cys Trp His His Pro Thr Ser Phe Glu Leu Cys Tyr
 65 70 75 80
 Pro Gly Thr Ser Ile Tyr Tyr Ala Ser Arg Gly Gly Pro Xaa Pro Asn
 85 90 95
 Ser Glu Xaa

<210> 337

<211> 96

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (1)

<223> Xaa equals any of the naturally occurring L-amino acids

<220>

<221> SITE

<222> (3)

<223> Xaa equals any of the naturally occurring L-amino acids

<400> 337

Xaa Asn Xaa Lys Ser Pro Leu Thr Ile Gly Asn Lys Ser Trp Ser Ser
 1 5 10 15
 Thr Ala Val Ala Ala Ala Leu Glu Leu Val Asp Pro Pro Gly Cys Arg
 20 25 30
 Asn Ser Ala Arg Asp Ser Pro Glu Leu Val His Leu Gly Lys Gly Arg
 35 40 45
 Pro Arg Lys Leu Met Thr Tyr Leu Phe Cys Ser Ser Ile Ser Leu Leu
 50 55 60
 Leu Leu Lys Val His Ser Ser Gly His Gln Asp Ile Arg Lys Ala Lys
 65 70 75 80
 Ser Lys Val Pro Arg Leu Leu Ile Ile Gln Cys Pro Gln Gln Arg Glu
 85 90 95

<210> 338

<211> 54

<212> PRT

<213> Homo sapiens

<400> 338

Gly Pro Glu Glu Asn Leu Ser Pro Ser Thr Pro Ser Gln Met Pro Thr
 1 5 10 15
 Ile Trp Val Lys Leu Cys Leu Leu Gln Val Cys His Gly Leu Phe Pro
 20 25 30
 Leu Leu Lys His Trp Ser Gln Pro Met Pro Leu Cys Val Thr Leu Ala
 35 40 45
 Pro Val Ser Tyr Trp Leu
 50

<210> 339

<211> 287

<212> PRT

<213> Homo sapiens

<400> 339

Pro Arg Val Arg Lys Glu Pro Glu Ala Met Gln Trp Leu Arg Val Arg
 5 10 15

Glu Ser Pro Gly Glu Ala Thr Gly His Arg Val Thr Met Gly Thr Ala
 20 25 30
 Ala Leu Gly Pro Val Trp Ala Ala Leu Leu Leu Phe Leu Leu Met Cys
 35 40 45
 Glu Ile Pro Met Val Glu Leu Thr Phe Asp Arg Ala Val Ala Ser Asp
 50 55 60
 Cys Gln Arg Cys Cys Asp Ser Glu Asp Pro Leu Asp Pro Ala His Val
 65 70 75 80
 Ser Ser Ala Ser Ser Ser Gly Arg Pro His Ala Leu Pro Glu Ile Arg
 85 90 95
 Pro Tyr Ile Asn Ile Thr Ile Leu Lys Gly Asp Lys Gly Asp Pro Gly
 100 105 110
 Pro Met Gly Leu Pro Gly Tyr Met Gly Arg Glu Gly Pro Gln Gly Glu
 115 120 125
 Pro Gly Pro Gln Gly Ser Lys Gly Asp Lys Gly Glu Met Gly Ser Pro
 130 135 140
 Gly Ala Pro Cys Gln Lys Arg Phe Phe Ala Phe Ser Val Gly Arg Lys
 145 150 155 160
 Thr Ala Leu His Ser Gly Glu Asp Phe Gln Thr Leu Leu Phe Glu Arg
 165 170 175
 Val Phe Val Asn Leu Asp Gly Cys Phe Asp Met Ala Thr Gly Gln Phe
 180 185 190
 Ala Ala Pro Leu Arg Gly Ile Tyr Phe Phe Ser Leu Asn Val His Ser
 195 200 205
 Trp Asn Tyr Lys Glu Thr Tyr Val His Ile Met His Asn Gln Lys Glu
 210 215 220
 Ala Val Ile Leu Tyr Ala Gln Pro Ser Glu Arg Ser Ile Met Gln Ser
 225 230 235 240
 Gln Ser Val Met Leu Asp Leu Ala Tyr Gly Asp Arg Val Trp Val Arg
 245 250 255
 Leu Phe Lys Arg Gln Arg Glu Asn Ala Ile Tyr Ser Asn Asp Phe Asp
 260 265 270
 Thr Tyr Ile Thr Phe Ser Gly His Leu Ile Lys Ala Glu Asp Asp
 275 280 285

<210> 340

<211> 339

<212> PRT

<213> Homo sapiens

<400> 340

Met Leu Tyr Pro Gly Ser Val Tyr Leu Leu Glu Lys Ala Leu Met Pro

1	5	10	15
Val Leu Leu Gln Gly Gln Ala Arg Leu Val Glu Glu Cys Asn Gly Arg	20	25	30
Arg Ala Lys Leu Leu Ala Cys Asp Gly Asn Glu Ile Asp Thr Met Phe	35	40	45
Val Asp Arg Arg Gly Thr Ala Glu Pro Gln Gly Gln Lys Leu Val Ile	50	55	60
Cys Cys Glu Gly Asn Ala Gly Phe Tyr Glu Val Gly Cys Val Ser Thr	65	70	75
Pro Leu Glu Ala Gly Tyr Ser Val Leu Gly Trp Asn His Pro Gly Phe	85	90	95
Ala Gly Ser Thr Gly Val Pro Phe Pro Gln Asn Glu Ala Asn Ala Met	100	105	110
Asp Val Val Val Gln Phe Ala Ile His Arg Leu Gly Phe Gln Pro Gln	115	120	125
Asp Ile Ile Ile Tyr Ala Trp Ser Ile Gly Gly Phe Thr Ala Thr Trp	130	135	140
Ala Ala Met Ser Tyr Pro Asp Val Ser Ala Met Ile Leu Asp Ala Ser	145	150	155
Phe Asp Asp Leu Val Pro Leu Ala Leu Lys Val Met Pro Asp Ser Trp	165	170	175
Arg Gly Leu Val Thr Arg Thr Val Arg Gln His Leu Asn Leu Asn Asn	180	185	190
Ala Glu Gln Leu Cys Arg Tyr Gln Gly Pro Val Leu Leu Ile Arg Arg	195	200	205
Thr Lys Asp Glu Ile Ile Thr Thr Thr Val Pro Glu Asp Ile Met Ser	210	215	220
Asn Arg Gly Asn Asp Leu Leu Leu Lys Leu Leu Gln His Arg Tyr Pro	225	230	235
Arg Val Met Ala Glu Glu Gly Leu Arg Val Val Arg Gln Trp Leu Glu	245	250	255
Ala Ser Ser Gln Leu Glu Glu Ala Ser Ile Tyr Ser Arg Trp Glu Val	260	265	270
Glu Glu Asp Trp Cys Leu Ser Val Leu Arg Ser Tyr Gln Ala Glu His	275	280	285
Gly Pro Asp Phe Pro Trp Ser Val Gly Glu Asp Met Ser Ala Asp Gly	290	295	300
Arg Arg Gln Leu Ala Leu Phe Leu Ala Arg Lys His Leu His Asn Phe	305	310	315
			320

Glu Ala Thr His Cys Thr Pro Leu Pro Ala Gln Asn Phe Gln Met Pro
325 330 335

Trp His Leu

<210> 341
<211> 127
<212> PRT
<213> Homo sapiens

<400> 341
Val Cys Pro Lys Trp Cys Arg Phe Leu Thr Met Leu Gly His Cys Cys
1 5 10 15
Tyr Phe Trp Gln Val Trp Pro Ala Ser Glu Ala Leu Ala Ala Gly Pro
20 25 30
Thr Pro Ser Thr Gly Ser Ser Ser Pro Ser Trp Lys Gln His Ile Gly
35 40 45
Thr Ser Leu Gln Lys Thr Arg Gly Ser Leu Pro Thr Thr Leu Thr
50 55 60
Ser Gly Ala Gly Gln Ser Thr Ser Thr Gly Lys Asn Pro Ala Ala Gly
65 70 75 80
Arg Ser Leu Glu Gly Ala Leu Pro Ala Gly Val Trp Pro Cys Phe Ala
85 90 95
Gln Ser Pro Cys Thr Gly Gly Gln Gln Thr Pro Ser Ser Thr Gly Leu
100 105 110
Arg Ser Cys Leu Val Arg Ser Pro Ala Thr Trp Trp Arg Thr Pro
115 120 125

<210> 342
<211> 554
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (16)
<223> Xaa equals any of the naturally occurring L-amino acids

<220>
<221> SITE
<222> (109)
<223> Xaa equals any of the naturally occurring L-amino acids

<400> 342
Trp Ile Pro Arg Ala Ala Gly Ile Arg His Glu Ile Tyr Arg Glu Xaa
1 5 10 15
Asp Ser Glu Arg Ala Pro Ala Ser Val Pro Glu Thr Pro Thr Ala Val
20 25 30

Thr Ala Pro His Ser Ser Ser Trp Asp Thr Tyr Tyr Gln Pro Arg Ala
35 40 45
Leu Glu Lys His Ala Asp Ser Ile Leu Ala Leu Ala Ser Val Phe Trp
50 55 60
Ser Ile Ser Tyr Tyr Ser Ser Pro Phe Ala Phe Phe Tyr Leu Tyr Arg
65 70 75 80
Lys Gly Tyr Leu Ser Leu Ser Lys Val Val Pro Phe Ser His Tyr Ala
85 90 95
Gly Thr Leu Leu Leu Leu Leu Ala Gly Val Ala Cys Xaa Arg Gly Ile
100 105 110
Gly Arg Trp Thr Asn Pro Gln Tyr Arg Gln Phe Ile Thr Ile Leu Glu
115 120 125
Ala Thr His Arg Asn Gln Ser Ser Glu Asn Lys Arg Gln Leu Ala Asn
130 135 140
Tyr Asn Phe Asp Phe Arg Ser Trp Pro Val Asp Phe His Trp Glu Glu
145 150 155 160
Pro Ser Ser Arg Lys Glu Ser Arg Gly Gly Pro Ser Arg Arg Gly Val
165 170 175
Ala Leu Leu Arg Pro Glu Pro Leu His Arg Gly Thr Ala Asp Thr Leu
180 185 190
Leu Asn Arg Val Lys Lys Leu Pro Cys Gln Ile Thr Ser Tyr Leu Val
195 200 205
Ala His Thr Leu Gly Arg Arg Met Leu Tyr Pro Gly Ser Val Tyr Leu
210 215 220
Leu Gln Lys Ala Leu Met Pro Val Leu Leu Gln Gly Gln Ala Arg Leu
225 230 235 240
Val Glu Glu Cys Asn Gly Arg Arg Ala Lys Leu Leu Ala Cys Asp Gly
245 250 255
Asn Glu Ile Asp Thr Met Phe Val Asp Arg Arg Gly Thr Ala Glu Pro
260 265 270
Gln Gly Gln Lys Leu Val Ile Cys Cys Glu Gly Asn Ala Gly Phe Tyr
275 280 285
Glu Val Gly Cys Val Ser Thr Pro Leu Glu Ala Gly Tyr Ser Val Leu
290 295 300
Gly Trp Asn His Pro Gly Phe Ala Gly Ser Thr Gly Val Pro Phe Pro
305 310 315 320
Gln Asn Glu Ala Asn Ala Met Asp Val Val Val Gln Phe Ala Ile His
325 330 335
Arg Leu Gly Phe Gln Pro Gln Asp Ile Ile Ile Tyr Ala Trp Ser Ile

340 345 350
 Gly Gly Phe Thr Ala Thr Trp Ala Ala Met Ser Tyr Pro Asp Val Ser
 355 360 365
 Ala Met Ile Leu Asp Ala Ser Phe Asp Asp Leu Val Pro Leu Ala Leu
 370 375 380
 Lys Val Met Pro Asp Ser Trp Arg Gly Leu Val Thr Arg Thr Val Arg
 385 390 395 400
 Gln His Leu Asn Leu Asn Asn Ala Glu Gln Leu Cys Arg Tyr Gln Gly
 405 410 415
 Pro Val Leu Leu Ile Arg Arg Thr Lys Asp Glu Ile Ile Thr Thr Thr
 420 425 430
 Val Pro Glu Asp Ile Met Ser Asn Arg Gly Asn Asp Leu Leu Leu Lys
 435 440 445
 Leu Leu Gln His Arg Tyr Pro Arg Val Met Ala Glu Glu Gly Leu Arg
 450 455 460
 Val Val Arg Gln Trp Leu Glu Ala Ser Ser Gln Leu Glu Glu Ala Ser
 465 470 475 480
 Ile Tyr Ser Arg Trp Glu Val Glu Glu Asp Trp Cys Leu Ser Val Leu
 485 490 495
 Arg Ser Tyr Gln Ala Glu His Gly Pro Asp Phe Pro Trp Ser Val Gly
 500 505 510
 Glu Asp Met Ser Ala Asp Gly Arg Arg Gln Leu Ala Leu Phe Leu Ala
 515 520 525
 Arg Lys His Leu His Asn Phe Glu Ala Thr His Cys Thr Pro Leu Pro
 530 535 540
 Ala Gln Asn Phe Gln Met Pro Trp His Leu
 545 550

<210> 343
 <211> 225
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (5)
 <223> Xaa equals any of the naturally occurring L-amino acids

<400> 343
 His Glu Arg Ala Xaa Gly Pro Ser Arg Gly His Gly Glu Leu Ser
 1 5 10 15
 Cys Val Leu Gly Pro Arg Leu Tyr Lys Ile Tyr Arg Glu Arg Asp Ser
 20 25 30

Glu Arg Ala Pro Ala Ser Val Pro Glu Thr Pro Thr Ala Val Thr Ala
 35 40 45
 Pro His Ser Ser Ser Trp Asp Thr Tyr Tyr Gln Pro Arg Ala Leu Glu
 50 55 60
 Lys His Ala Asp Ser Ile Leu Ala Leu Ala Ser Val Phe Trp Ser Ile
 65 70 75 80
 Ser Tyr Tyr Ser Ser Pro Phe Ala Phe Phe Tyr Leu Tyr Arg Lys Gly
 85 90 95
 Tyr Leu Ser Leu Ser Lys Val Val Pro Phe Ser His Tyr Ala Gly Thr
 100 105 110
 Leu Leu Leu Leu Leu Ala Gly Val Ala Cys Ser Glu Ala Leu Ala Ala
 115 120 125
 Gly Pro Thr Pro Ser Thr Gly Ser Ser Ser Pro Ser Trp Lys Gln His
 130 135 140
 Ile Gly Thr Ser Leu Gln Lys Thr Arg Gly Ser Leu Pro Thr Thr Thr
 145 150 155 160
 Leu Thr Ser Gly Ala Gly Gln Ser Thr Ser Thr Gly Lys Asn Pro Ala
 165 170 175
 Ala Gly Arg Ser Leu Glu Gly Ala Leu Pro Ala Gly Val Trp Pro Cys
 180 185 190
 Phe Ala Gln Ser Pro Cys Thr Gly Gly Gln Gln Thr Pro Ser Ser Thr
 195 200 205
 Gly Leu Arg Ser Cys Leu Val Arg Ser Pro Ala Thr Trp Trp Arg Thr
 210 215 220
 Pro
 225

<210> 344
 <211> 299
 <212> PRT
 <213> Homo sapiens

<400> 344
 Met Phe Lys Arg His Gln Arg Leu Lys Lys Asp Ser Thr Gln Ala Glu
 1 5 10 15
 Glu Asp Leu Ser Glu Gln Glu Gln Asn Gln Leu Asn Val Leu Lys Lys
 20 25 30
 His Gly Tyr Val Val Gly Arg Val Gly Arg Thr Phe Leu Tyr Ser Glu
 35 40 45
 Glu Gln Lys Asp Asn Ile Pro Phe Glu Phe Asp Ala Asp Ser Leu Ala
 50 55 60
 Phe Asp Met Glu Asn Asp Pro Val Met Gly Thr His Lys Ser Thr Lys

65		70		75		80
Gln Val Glu Leu Thr	Ala Gln Asp Val	Lys Asp Ala His Trp Phe Tyr				
	85	90				95
Asp Thr Pro Gly Ile Thr Lys Glu Asn Cys Ile Leu Asn Leu Leu Thr						
	100	105				110
Glu Lys Glu Val Asn Ile Val Leu Pro Thr Gln Ser Ile Val Pro Arg						
	115	120				125
Thr Phe Val Leu Lys Pro Gly Met Val Leu Phe Leu Gly Ala Ile Gly						
	130	135				140
Arg Ile Asp Phe Leu Gln Gly Asn Gln Ser Ala Trp Phe Thr Val Val						
	145	150				155
Ala Ser Asn Ile Leu Pro Val His Ile Thr Ser Leu Asp Arg Ala Asp						
	165	170				175
Ala Leu Tyr Gln Lys His Ala Gly His Thr Leu Leu Gln Ile Pro Met						
	180	185				190
Gly Gly Lys Glu Arg Met Ala Gly Phe Pro Pro Leu Val Ala Glu Asp						
	195	200				205
Ile Met Leu Lys Glu Gly Leu Gly Ala Ser Glu Ala Val Ala Asp Ile						
	210	215				220
Lys Phe Ser Ser Ala Gly Trp Val Ser Val Thr Pro Asn Phe Lys Asp						
	225	230				235
Arg Leu His Leu Arg Gly Tyr Thr Pro Glu Gly Thr Val Leu Thr Val						
	245	250				255
Arg Pro Pro Leu Leu Pro Tyr Ile Val Asn Ile Lys Gly Gln Arg Ile						
	260	265				270
Lys Lys Ser Val Ala Tyr Lys Thr Lys Lys Pro Pro Ser Leu Met Tyr						
	275	280				285
Asn Val Arg Lys Lys Lys Gly Lys Ile Asn Val						
	290	295				

<210> 345

<211> 314

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (147)

<223> Xaa equals any of the naturally occurring L-amino acids

<220>

<221> SITE

<222> (211)

<223> Xaa equals any of the naturally occurring L-amino acids

<400> 345
 Met Leu Pro Ala Arg Leu Pro Phe Arg Leu Leu Ser Leu Phe Leu Arg
 1 5 10 15
 Gly Ser Ala Pro Thr Ala Ala Arg His Gly Leu Arg Glu Pro Leu Leu
 20 25 30
 Glu Arg Arg Cys Ala Ala Ala Ser Ser Phe Gln His Ser Ser Ser Leu
 35 40 45
 Gly Arg Glu Leu Pro Tyr Asp Pro Val Asp Thr Glu Gly Phe Gly Glu
 50 55 60
 Gly Gly Asp Met Gln Glu Arg Phe Leu Phe Pro Glu Tyr Ile Leu Asp
 65 70 75 80
 Pro Glu Pro Gln Pro Thr Arg Glu Lys Gln Leu Gln Glu Leu Gln Gln
 85 90 95
 Gln Gln Glu Glu Glu Glu Arg Gln Arg Gln Gln Arg Arg Glu Glu Arg
 100 105 110
 Arg Gln Gln Asn Leu Arg Ala Arg Ser Arg Glu His Pro Val Val Gly
 115 120 125
 His Pro Asp Pro Ala Leu Pro Pro Ser Gly Val Asn Cys Ser Gly Cys
 130 135 140
 Gly Ala Xaa Leu His Cys Gln Asp Ala Gly Val Pro Gly Tyr Leu Pro
 145 150 155 160
 Arg Glu Lys Phe Leu Arg Thr Ala Glu Ala Asp Gly Gly Leu Ala Arg
 165 170 175
 Thr Val Cys Gln Arg Cys Trp Leu Leu Ser His His Arg Arg Ala Leu
 180 185 190
 Arg Leu Gln Val Ser Arg Glu Gln Tyr Leu Glu Leu Val Ser Ala Ala
 195 200 205
 Leu Arg Xaa Pro Gly Pro Ser Leu Val Leu Tyr Met Val Asp Leu Leu
 210 215 220
 Asp Leu Pro Asp Ala Leu Leu Pro Asp Leu Pro Ala Leu Val Gly Pro
 225 230 235 240
 Lys Gln Leu Ile Val Leu Gly Asn Lys Val Asp Leu Leu Pro Gln Asp
 245 250 255
 Ala Pro Gly Tyr Arg Gln Arg Leu Arg Glu Arg Leu Trp Glu Asp Cys
 260 265 270
 Ala Arg Ala Gly Leu Leu Leu Ala Pro Gly Thr Lys Gly His Ser Ala
 275 280 285
 Pro Ser Arg Thr Ser His Arg Thr Gly Arg Ile Arg Ile Arg Arg Thr
 290 295 300

Gly Pro Ala Gln Trp Ser Gly Thr Cys Gly
305 310

<210> 346
<211> 380
<212> PRT
<213> Homo sapiens

<400> 346
Pro Ser Phe Arg Arg Glu Arg Val Glu Thr Gly Gly Gly Gly Pro Val
1 5 10 15
Thr His Gly Thr Glu Gly Pro Phe Leu Pro Leu Pro Gly Gly Thr Arg
20 25 30
Met Asn Met Thr Gln Ala Arg Val Leu Val Ala Ala Val Val Gly Leu
35 40 45
Val Ala Val Leu Leu Tyr Ala Ser Ile His Lys Ile Glu Glu Gly His
50 55 60
Leu Ala Val Tyr Tyr Arg Gly Gly Ala Leu Leu Thr Ser Pro Ser Gly
65 70 75 80
Pro Gly Tyr His Ile Met Leu Pro Phe Ile Thr Thr Phe Arg Ser Val
85 90 95
Gln Thr Thr Leu Gln Thr Asp Glu Val Lys Asn Val Pro Cys Gly Thr
100 105 110
Ser Gly Gly Val Met Ile Tyr Ile Asp Arg Ile Glu Val Val Asn Met
115 120 125
Leu Ala Pro Tyr Ala Val Phe Asp Ile Val Arg Asn Tyr Thr Ala Asp
130 135 140
Tyr Asp Lys Thr Leu Ile Phe Asn Lys Ile His His Glu Leu Asn Gln
145 150 155 160
Phe Cys Ser Ala His Thr Leu Gln Glu Val Tyr Ile Glu Leu Phe Asp
165 170 175
Gln Ile Asp Glu Asn Leu Lys Gln Ala Leu Gln Lys Asp Leu Asn Leu
180 185 190
Met Ala Pro Gly Leu Thr Ile Gln Ala Val Arg Val Thr Lys Pro Lys
195 200 205
Ile Pro Glu Ala Ile Arg Arg Asn Phe Glu Leu Met Glu Ala Glu Lys
210 215 220
Thr Lys Leu Leu Ile Ala Ala Gln Lys Gln Lys Val Val Glu Lys Glu
225 230 235 240
Ala Glu Thr Glu Arg Lys Lys Ala Val Ile Glu Ala Glu Lys Ile Ala
245 250 255
Gln Val Ala Lys Ile Arg Phe Gln Gln Lys Val Met Glu Lys Glu Thr

260 265 270
 Glu Lys Arg Ile Ser Glu Ile Glu Asp Ala Ala Phe Leu Ala Arg Glu
 275 280 285
 Lys Ala Lys Ala Asp Ala Glu Tyr Tyr Ala Ala His Lys Tyr Ala Thr
 290 295 300
 Ser Asn Lys His Lys Leu Thr Pro Glu Tyr Leu Glu Leu Lys Lys Tyr
 305 310 315 320
 Gln Ala Ile Ala Ser Asn Ser Lys Ile Tyr Phe Gly Ser Asn Ile Pro
 325 330 335
 Asn Met Phe Val Asp Ser Ser Cys Ala Leu Lys Tyr Ser Asp Ile Arg
 340 345 350
 Thr Gly Arg Glu Ser Ser Leu Pro Ser Lys Glu Ala Leu Glu Pro Ser
 355 360 365
 Gly Glu Asn Val Ile Gln Asn Lys Glu Ser Thr Gly
 370 375 380

 <210> 347
 <211> 422
 <212> PRT
 <213> Homo sapiens

 <220>
 <221> SITE
 <222> (328)
 <223> Xaa equals any of the naturally occurring L-amino acids

 <400> 347
 Trp Ser Thr Gly Asn Ala Ser Trp Glu Lys Lys Asp Asn Phe Ile Leu
 1 5 10 15
 Ser Ala Asp Phe Glu Met Met Gly Leu Gly Asn Gly Arg Arg Ser Met
 20 25 30
 Lys Ser Pro Pro Leu Val Leu Ala Ala Leu Val Ala Cys Ile Ile Val
 35 40 45
 Leu Gly Phe Asn Tyr Trp Ile Ala Ser Ser Arg Ser Val Asp Leu Gln
 50 55 60
 Thr Arg Ile Met Glu Leu Glu Gly Arg Val Arg Arg Arg Ala Ala Glu
 65 70 75 80
 Arg Gly Ala Val Glu Leu Lys Lys Asn Glu Phe Gln Gly Glu Leu Glu
 85 90 95
 Lys Gln Arg Glu Gln Leu Asp Lys Ile Gln Ser Ser His Asn Phe Gln
 100 105 110
 Leu Glu Ser Val Asn Lys Leu Tyr Gln Asp Glu Lys Ala Val Leu Val
 115 120 125

Asn Asn Ile Thr Thr Gly Glu Arg Leu Ile Arg Val Leu Gln Asp Gln
 130 135 140
 Leu Lys Thr Leu Gln Arg Asn Tyr Gly Arg Leu Gln Gln Asp Val Leu
 145 150 155 160
 Gln Phe Gln Lys Asn Gln Thr Asn Leu Glu Arg Lys Phe Ser Tyr Asp
 165 170 175
 Leu Ser Gln Cys Ile Asn Gln Met Lys Glu Val Lys Glu Gln Cys Glu
 180 185 190
 Glu Arg Ile Glu Glu Val Thr Lys Lys Gly Asn Glu Ala Val Ala Ser
 195 200 205
 Arg Asp Leu Ser Glu Asn Asn Asp Gln Arg Gln Gln Leu Gln Ala Leu
 210 215 220
 Ser Glu Pro Gln Pro Arg Leu Gln Ala Ala Gly Leu Pro His Thr Glu
 225 230 235 240
 Val Pro Gln Gly Lys Gly Asn Val Leu Gly Asn Ser Lys Ser Gln Thr
 245 250 255
 Pro Ala Pro Ser Ser Glu Val Val Leu Asp Ser Lys Arg Gln Val Glu
 260 265 270
 Lys Glu Glu Thr Asn Glu Ile Gln Val Val Asn Glu Glu Pro Gln Arg
 275 280 285
 Asp Arg Leu Pro Gln Glu Pro Gly Arg Glu Gln Val Val Glu Asp Arg
 290 295 300
 Pro Val Gly Gly Arg Gly Phe Gly Gly Ala Gly Glu Leu Gly Gln Thr
 305 310 315 320
 Pro Gln Val Gln Ala Ala Leu Xaa Val Ser Gln Glu Asn Pro Glu Met
 325 330 335
 Glu Gly Pro Glu Arg Asp Gln Leu Val Ile Pro Asp Gly Gln Glu Glu
 340 345 350
 Glu Gln Glu Ala Ala Gly Glu Gly Arg Asn Gln Gln Lys Leu Arg Gly
 355 360 365
 Glu Asp Asp Tyr Asn Met Asp Glu Asn Glu Ala Glu Ser Glu Thr Asp
 370 375 380
 Lys Gln Ala Ala Leu Ala Gly Asn Asp Arg Asn Ile Asp Val Phe Asn
 385 390 395 400
 Val Glu Asp Gln Lys Arg Asp Thr Ile Asn Leu Leu Asp Gln Arg Glu
 405 410 415
 Lys Arg Asn His Thr Leu
 420

<210> 348

<211> 14
<212> PRT
<213> Homo sapiens

<400> 348
Ser Leu His Arg Phe Val Leu Ser Gln Ala Lys Asp Glu Leu
1 5 10

<210> 349
<211> 19
<212> PRT
<213> Homo sapiens

<400> 349
Phe Ile Lys Phe Phe Ala Pro Trp Cys Gly His Cys Lys Ala Leu Ala
1 5 10 15

Pro Thr Trp

<210> 350
<211> 19
<212> PRT
<213> Homo sapiens

<400> 350
Phe Ile Lys Phe Tyr Ala Pro Trp Cys Gly His Cys Lys Thr Leu Ala
1 5 10 15

Pro Thr Trp

<210> 351
<211> 363
<212> PRT
<213> Homo sapiens

<220>
<221> SITE
<222> (42)
<223> Xaa equals any of the naturally occurring L-amino acids

<400> 351
Arg Arg Gly Arg Gly Val Pro Gly Pro Arg Gly Arg Arg Arg Leu Trp
1 5 10 15

Ser Ala Ala Cys Gly His Cys Gln Arg Leu Gln Pro Thr Trp Asn Asp
20 25 30

Leu Gly Asp Lys Tyr Asn Ser Met Glu Xaa Ala Lys Val Tyr Val Ala
35 40 45

Lys Val Asp Cys Thr Ala His Ser Asp Val Cys Ser Ala Gln Gly Val
50 55 60

Arg Gly Tyr Pro Thr Leu Lys Leu Phe Lys Pro Gly Gln Glu Ala Val

65		70		75		80
Lys Tyr Gln Gly Pro Arg Asp Phe Gln Thr Leu Glu Asn Trp Met Leu						
	85			90		95
Gln Thr Leu Asn Glu Glu Pro Val Thr Pro Glu Pro Glu Val Glu Pro						
	100		105			110
Pro Ser Ala Pro Glu Leu Lys Gln Gly Leu Tyr Glu Leu Ser Ala Ser						
	115		120			125
Asn Phe Glu Leu His Val Ala Gln Gly Asp His Phe Ile Lys Phe Phe						
	130		135			140
Ala Pro Trp Cys Gly His Cys Lys Ala Leu Ala Pro Thr Trp Glu Gln						
	145		150		155	160
Leu Ala Leu Gly Leu Glu His Ser Glu Thr Val Lys Ile Gly Lys Val						
	165		170			175
Asp Cys Thr Gln His Tyr Glu Leu Cys Ser Gly Asn Gln Val Arg Gly						
	180		185			190
Tyr Pro Thr Leu Leu Trp Phe Arg Asp Gly Lys Lys Val Asp Gln Tyr						
	195		200			205
Lys Gly Lys Arg Asp Leu Glu Ser Leu Arg Glu Tyr Val Glu Ser Gln						
	210		215			220
Leu Gln Arg Thr Glu Thr Gly Ala Thr Glu Thr Val Thr Pro Ser Glu						
	225		230		235	240
Ala Pro Val Leu Ala Ala Glu Pro Glu Ala Asp Lys Gly Thr Val Leu						
	245			250		255
Ala Leu Thr Glu Asn Asn Phe Asp Asp Thr Ile Ala Glu Gly Ile Thr						
	260		265			270
Phe Ile Lys Phe Tyr Ala Pro Trp Cys Gly His Cys Lys Thr Leu Ala						
	275		280			285
Pro Thr Trp Glu Glu Leu Ser Lys Lys Glu Phe Pro Gly Leu Ala Gly						
	290		295			300
Val Lys Ile Ala Glu Val Asp Cys Thr Ala Glu Arg Asn Ile Cys Ser						
	305		310		315	320
Lys Tyr Ser Val Arg Gly Tyr Pro Thr Leu Leu Leu Phe Arg Gly Gly						
	325			330		335
Lys Lys Val Ser Glu His Ser Gly Gly Arg Asp Leu Asp Ser Leu His						
	340			345		350
Arg Phe Val Leu Ser Gln Ala Lys Asp Glu Leu						
	355			360		

<210> 352

<211> 93

<212> PRT

<213> Homo sapiens

<400> 352

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Met Arg Pro Gln Gly Pro Ala Ala Ser Pro Gln Arg Leu Arg Gly Leu
 1           5           10           15
Leu Leu Leu Leu Leu Leu Gln Leu Pro Ala Pro Ser Ser Ala Ser Glu
 20           25           30
Ile Pro Lys Gly Lys Gln Lys Ala His Ser Gly Arg Gly Arg Trp Trp
 35           40           45
Thr Cys Ile Met Glu Cys Ala Tyr Lys Gly Gln Gln Glu Cys Leu Val
 50           55           60
Glu Thr Gly Ala Leu Gly Pro Met Ala Phe Arg Val His Leu Gly Ser
 65           70           75           80
Gln Val Gly Met Asp Ser Lys Glu Lys Arg Gly Asn Val
           85           90

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<210> 353

<211> 273

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (210)

<223> Xaa equals any of the naturally occurring L-amino acids

<400> 353

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Glu Thr Arg Val Lys Thr Ser Leu Glu Leu Arg Thr Gln Leu Glu
 1           5           10           15
Pro Thr Gly Thr Val Gly Asn Thr Ile Met Thr Ser Gln Pro Val Pro
 20           25           30
Asn Glu Thr Ile Ile Val Leu Pro Ser Asn Val Ile Asn Phe Ser Gln
 35           40           45
Ala Glu Lys Pro Glu Pro Thr Asn Gln Gly Gln Asp Ser Leu Lys Lys
 50           55           60
His Leu His Ala Glu Ile Lys Val Ile Gly Thr Ile Gln Ile Leu Cys
 65           70           75           80
Gly Met Met Val Leu Ser Leu Gly Ile Ile Leu Ala Ser Ala Ser Phe
 85           90           95
Ser Pro Asn Phe Thr Glp Val Thr Ser Thr Leu Leu Asn Ser Ala Tyr
 100          105          110
Pro Phe Ile Gly Pro Phe Phe Phe Ile Ile Ser Gly Ser Leu Ser Ile
 115          120          125
Ala Thr Glu Lys Arg Leu Thr Lys Leu Leu Val His Ser Ser Leu Val

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130 135 140
 Gly Ser Ile Leu Ser Ala Leu Ser Ala Leu Val Gly Phe Ile Ile Leu
 145 150 155 160
 Ser Val Lys Gln Ala Thr Leu Asn Pro Ala Ser Leu Gln Cys Glu Leu
 165 170 175
 Asp Lys Asn Asn Ile Pro Thr Arg Ser Tyr Val Ser Tyr Phe Tyr His
 180 185 190
 Asp Ser Leu Tyr Thr Thr Asp Cys Tyr Thr Ala Lys Ala Ser Leu Ala
 195 200 205
 Gly Xaa Leu Ser Leu Met Leu Ile Cys Thr Leu Leu Glu Phe Cys Leu
 210 215 220
 Ala Val Leu Thr Ala Val Leu Arg Trp Lys Gln Ala Tyr Ser Asp Phe
 225 230 235 240
 Pro Gly Ser Val Leu Phe Leu Pro His Ser Tyr Ile Gly Asn Ser Gly
 245 250 255
 Met Ser Ser Lys Met Thr His Asp Cys Gly Tyr Glu Glu Leu Leu Thr
 260 265 270
 Ser

<210> 354

<211> 192

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (129)

<223> Xaa equals any of the naturally occurring L-amino acids

<400> 354

Met Met Val Leu Ser Leu Gly Ile Ile Leu Ala Ser Ala Ser Phe Ser
 1 5 10 15
 Pro Asn Phe Thr Gln Val Thr Ser Thr Leu Leu Asn Ser Ala Tyr Pro
 20 25 30
 Phe Ile Gly Pro Phe Phe Phe Ile Ile Ser Gly Ser Leu Ser Ile Ala
 35 40 45
 Thr Glu Lys Arg Leu Thr Lys Leu Leu Val His Ser Ser Leu Val Gly
 50 55 60
 Ser Ile Leu Ser Ala Leu Ser Ala Leu Val Gly Phe Ile Ile Leu Ser
 65 70 75 80
 Val Lys Gln Ala Thr Leu Asn Pro Ala Ser Leu Gln Cys Glu Leu Asp
 85 90 95

Lys Asn Asn Ile Pro Thr Arg Ser Tyr Val Ser Tyr Phe Tyr His Asp
 100 105 110
 Ser Leu Tyr Thr Thr Asp Cys Tyr Thr Ala Lys Ala Ser Leu Ala Gly
 115 120 125
 Xaa Leu Ser Leu Met Leu Ile Cys Thr Leu Leu Glu Phe Cys Leu Ala
 130 135 140
 Val Leu Thr Ala Val Leu Arg Trp Lys Gln Ala Tyr Ser Asp Phe Pro
 145 150 155 160
 Gly Ser Val Leu Phe Leu Pro His Ser Tyr Ile Gly Asn Ser Gly Met
 165 170 175
 Ser Ser Lys Met Thr His Asp Cys Gly Tyr Glu Glu Leu Leu Thr Ser
 180 185 190

<210> 355

<211> 204

<212> PRT

<213> Homo sapiens

<220>

<221> SITE

<222> (119)

<223> Xaa equals any of the naturally occurring L-amino acids

<400> 355

Gly Ala Ser Cys Glu Gly Gly Gly Ala Ala Ala Arg Ala Ala Leu Gly
 1 5 10 15
 Val His Arg Ser Gln Lys Ala Leu Leu Val Phe Arg Arg Thr Leu Ser
 20 25 30
 Asn Leu Leu Tyr Met Pro Leu Leu Arg Gly Leu Leu Trp Leu Gln Val
 35 40 45
 Leu Cys Ala Gly Pro Leu His Thr Glu Ala Val Val Leu Leu Val Pro
 50 55 60
 Ser Asp Asp Gly Arg Ala Phe Leu Leu Arg Ser Arg Leu Leu His Pro
 65 70 75 80
 Glu Ala His Val Pro Pro Ala Ala Asp Arg Gly Ala Ser Leu Gln Cys
 85 90 95
 Val Leu His Gln Ala Ala Pro Lys Ser Arg Pro Arg Ser Pro Ala Ala
 100 105 110
 Gly Ala Ala Leu Leu His Xaa Pro Arg Arg Thr Gly Asp Glu Pro Cys
 115 120 125
 Arg Glu Phe His Gly Asn Gly Phe Pro Gly Pro Thr Gln Leu Thr Pro
 130 135 140

Gly Glu Cys Gly Leu Pro Ala Pro Ser Ser Leu Leu Gln His Ala Ser
 145 150 155 160
 Ala Pro Val Arg Thr Gly Ser Glu Gly Gln Val Val Gly Cys Pro Arg
 165 170 175
 Ala Arg Gly Glu Thr Gly Glu Gly Leu Ser Leu Ala Phe Leu Ser Ser
 180 185 190
 Leu Met Phe Thr Ser Arg Asn Gly Leu Val Gly Cys
 195 200

<210> 356
 <211> 72
 <212> PRT
 <213> Homo sapiens

<400> 356
 Met Gly Ser Ala Ala Leu Glu Ile Leu Gly Leu Val Leu Cys Leu Val
 1 5 10 15
 Gly Trp Gly Gly Leu Ile Leu Ala Cys Gly Leu Pro Met Trp Gln Val
 20 25 30
 Thr Ala Phe Leu Asp His Asn Ile Val Thr Ala Gln Thr Thr Trp Lys
 35 40 45
 Gly Leu Trp Met Ser Cys Val Val Gln Ser Thr Gly Thr Cys Ser Ala
 50 55 60
 Lys Cys Thr Thr Arg Cys Trp Leu
 65 70

<210> 357
 <211> 115
 <212> PRT
 <213> Homo sapiens

<400> 357
 Leu Lys Arg Ala Pro Pro Gly Pro Ala Leu Ala Lys Gly Leu Leu Gln
 1 5 10 15
 Pro Ser Ser Thr Phe Gln Ala Leu Glu Thr Asn Ile Gly Asp Gln Val
 20 25 30
 Arg Arg His Ser Thr Ala Val Val Ile Arg Glu Met Thr Ser Tyr Ile
 35 40 45
 Leu Ile Ser Phe Val Leu Leu Ile Gly Val Gly Cys Ile Glu Lys Asp
 50 55 60
 Gln Ser Cys Pro Val Phe Gly Gly Arg Lys Arg Leu His Leu Leu Phe
 65 70 75 80
 Val Gly Gly Gln Leu Arg Gln Val Arg Met Leu Arg Gly Glu Leu Ser
 85 90 95

Cys Ala Cys Tyr Arg Pro His Val Gln Ala Leu Gln Leu Gly Gly Cys
 100 105 110

Thr Cys Phe
 115

<210> 358
 <211> 88
 <212> PRT
 <213> Homo sapiens

<400> 358
 Val Ile Lys Leu Ile Cys Pro Ala Ala Phe Pro Val Tyr Phe Gln Asp
 1 5 10 15
 Met Ala Arg Gly Cys Val Cys Ser Leu Cys Ala Ser Val Cys Ile Phe
 20 25 30
 Leu Ser Ser Leu Phe Pro Leu Leu Pro Ser Val His Ser Val Asn Ile
 35 40 45
 Ile Ser Cys Leu Leu Leu Ser Lys Cys Phe Glu Gly Leu Glu Leu Met
 50 55 60
 Cys Glu His Leu Tyr Gln Leu Ser Gln Leu His Val Leu His His Ile
 65 70 75 80
 Phe Ser Tyr Leu Leu Cys Thr Pro
 85

<210> 359
 <211> 716
 <212> PRT
 <213> Homo sapiens

<220>
 <221> SITE
 <222> (2)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (373)
 <223> Xaa equals any of the naturally occurring L-amino acids

<220>
 <221> SITE
 <222> (705)
 <223> Xaa equals any of the naturally occurring L-amino acids

<400> 359
 Tyr Xaa Ile Pro Gly Ser Thr His Ala Ser Gly Arg Gln Arg Gly Ser
 1 5 10 15
 Gly Arg Gly Glu Asp Asp Ser Gly Pro Pro Pro Ser Thr Val Ile Asn
 20 25 30

Gln Asn Glu Thr Phe Ala Asn Ile Ile Phe Lys Pro Thr Val Val Gln
 35 40 45
 Gln Ala Arg Ile Ala Gln Asn Gly Ile Leu Gly Asp Phe Ile Ile Arg
 50 55 60
 Tyr Asp Val Asn Arg Glu Gln Ser Ile Gly Asp Ile Gln Val Leu Asn
 65 70 75 80
 Gly Tyr Phe Val His Tyr Phe Ala Pro Lys Asp Leu Pro Pro Leu Pro
 85 90 95
 Lys Asn Val Val Phe Val Leu Asp Ser Ser Ala Ser Met Val Gly Thr
 100 105 110
 Lys Leu Arg Gln Thr Lys Asp Ala Leu Phe Thr Ile Leu His Asp Leu
 115 120 125
 Arg Pro Gln Asp Arg Phe Ser Ile Ile Gly Phe Ser Asn Arg Ile Lys
 130 135 140
 Val Trp Lys Asp His Leu Ile Ser Val Thr Pro Asp Ser Ile Arg Asp
 145 150 155 160
 Gly Lys Val Tyr Ile His His Met Ser Pro Thr Gly Gly Thr Asp Ile
 165 170 175
 Asn Gly Val Leu Gln Arg Ala Ile Arg Leu Leu Asn Lys Tyr Val Ala
 180 185 190
 His Ser Gly Ile Gly Asp Arg Ser Val Ser Leu Ile Val Phe Leu Thr
 195 200 205
 Asp Gly Lys Pro Thr Val Gly Glu Thr His Thr Leu Lys Ile Leu Asn
 210 215 220
 Asn Thr Arg Glu Ala Ala Arg Gly Gln Val Cys Ile Phe Thr Ile Gly
 225 230 235 240
 Ile Gly Asn Asp Val Asp Phe Arg Leu Leu Glu Lys Leu Ser Leu Glu
 245 250 255
 Asn Cys Gly Leu Thr Arg Arg Val His Glu Glu Glu Asp Ala Gly Ser
 260 265 270
 Gln Leu Ile Gly Phe Tyr Asp Glu Ile Arg Thr Pro Leu Leu Ser Asp
 275 280 285
 Ile Arg Ile Asp Tyr Pro Pro Ser Ser Val Val Gln Ala Thr Lys Thr
 290 295 300
 Leu Phe Pro Asn Tyr Phe Asn Gly Ser Glu Ile Ile Ile Ala Gly Lys
 305 310 315 320
 Leu Val Asp Arg Lys Leu Asp His Leu His Val Glu Val Thr Ala Ser
 325 330 335
 Asn Ser Lys Lys Phe Ile Ile Leu Lys Thr Asp Val Pro Val Arg Pro

340	345	350
Gln Lys Ala Gly Lys Asp Val Thr Gly Ser Pro Arg Pro Gly Gly Asp		
355	360	365
Gly Glu Gly Asp Xaa Asn His Ile Glu Arg Leu Trp Ser Tyr Leu Thr		
370	375	380
Thr Lys Glu Leu Leu Ser Ser Trp Leu Gln Ser Asp Asp Glu Pro Glu		
385	390	395
Lys Glu Arg Leu Arg Gln Arg Ala Gln Ala Leu Ala Val Ser Tyr Arg		
405	410	415
Phe Leu Thr Pro Phe Thr Ser Met Lys Leu Arg Gly Pro Val Pro Arg		
420	425	430
Met Asp Gly Leu Glu Glu Ala His Gly Met Ser Ala Ala Met Gly Pro		
435	440	445
Glu Pro Val Val Gln Ser Val Arg Gly Ala Gly Thr Gln Pro Gly Pro		
450	455	460
Leu Leu Lys Lys Pro Tyr Gln Pro Arg Ile Lys Ile Ser Lys Thr Ser		
465	470	475
Val Asp Gly Asp Pro His Phe Val Val Asp Phe Pro Leu Ser Arg Leu		
485	490	495
Thr Val Cys Phe Asn Ile Asp Gly Gln Pro Gly Asp Ile Leu Arg Leu		
500	505	510
Val Ser Asp His Arg Asp Ser Gly Val Thr Val Asn Gly Glu Leu Ile		
515	520	525
Gly Ala Pro Ala Pro Pro Asn Gly His Lys Lys Gln Arg Thr Tyr Leu		
530	535	540
Arg Thr Ile Thr Ile Leu Ile Asn Lys Pro Glu Arg Ser Tyr Leu Glu		
545	550	555
Ile Thr Pro Ser Arg Val Ile Leu Asp Gly Gly Asp Arg Leu Val Leu		
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Pro Cys Asn Gln Ser Val Val Val Gly Ser Trp Gly Leu Glu Val Ser		
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Val Ser Ala Asn Ala Asn Val Thr Val Thr Ile Gln Gly Ser Ile Ala		
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Phe Val Ile Leu Ile His Leu Tyr Lys Lys Pro Ala Pro Phe Gln Arg		
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Cys His Gly Leu Leu Gly Gln Phe Leu Asn Gln Asp Ala Arg Leu Thr		
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 660 665 670
 Gln Val Gly Glu Gly Pro Glu Ala Val Leu Thr Val Lys Gly His Gln
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 <212> PRT
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 Leu Ala Asn Gly His Ala Pro Tyr Ser Arg Thr Leu Ser His Ile Ser
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 <211> 155
 <212> PRT
 <213> Homo sapiens

<400> 362
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 Ser Leu His Trp Phe Ile Phe Leu Leu Asn Leu Pro Val Ala Thr Trp
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Asn Ile Tyr Arg Tyr Ile Met Val Pro Ser Gly Asn Met Gly Val Phe
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 Ser Ser Glu Ser Lys Ser Ser Leu
 50 55

Applicant's or agent's file reference number	P2031PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>280</u> , line <u>N/A</u>		REC'D 18 AUG 1999
		WIPO PCT
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>		
Name of depositary institution American Type Culture Collection		
Address of depositary institution (including postal code and country) 10801 University Boulevard Manassas, Virginia 20110-2209 United States of America		
Date of deposit July 27, 1998	Accession Number 203069	
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>		
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)		
Europe In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28(4) EPC).		
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)		
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g. "Accession Number of Deposit")		
For receiving Office use only		For International Bureau use only
<input checked="" type="checkbox"/> This sheet was received with the international application		<input checked="" type="checkbox"/> This sheet was received by the International Bureau on:
Authorized officer YVES E. SIMONE PCT International Division		18 AUGUST 1999 Authorized officer P. G. J. J.

CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

The applicant hereby requests that the application has been laid open to public inspection (by the Norwegian Patent Office), or has been finally decided upon by the Norwegian Patent Office without having been laid open inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Norwegian Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Norwegian Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on the list of recognized experts drawn up by the Norwegian Patent Office or any person approved by the applicant in the individual case.

AUSTRALIA

The applicant hereby gives notice that the furnishing of a sample of a microorganism shall only be effected prior to the grant of a patent, or prior to the lapsing, refusal or withdrawal of the application, to a person who is a skilled addressee without an interest in the invention (Regulation 3.25(3) of the Australian Patents Regulations).

FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

The applicant hereby requests that, until the application has been laid open to public inspection (by the Danish Patent Office), or has been finally decided upon by the Danish Patent office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Danish Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Danish Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Danish Patent Office or any person by the applicant in the individual case.

SWEDEN

The applicant hereby requests that, until the application has been laid open to public inspection (by the Swedish Patent Office), or has been finally decided upon by the Swedish Patent Office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the International Bureau before the expiration of 16 months from the priority date (preferably on the Form PCT/RO/134 reproduced in annex Z of Volume I of the PCT Applicant's Guide). If such a request has been filed by the applicant any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Swedish Patent Office or any person approved by a applicant in the individual case.

NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

Applicant's or agent's file reference number	PZ031PCT	International application:	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>243</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	
Date of deposit June 11, 1998	Accession Number 209965
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States) Europe In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28(4) EPC).	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposit")	
For receiving Office use only <input checked="" type="checkbox"/> This sheet was received with the international application Authorized officer Yvette S. Smith PCT International Division	For International Bureau use only <input type="checkbox"/> This sheet was received by the International Bureau on: Authorized officer

CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

The applicant hereby requests that the application has been laid open to public inspection (by the Norwegian Patent Office), or has been finally decided upon by the Norwegian Patent Office without having been laid open inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Norwegian Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Norwegian Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on the list of recognized experts drawn up by the Norwegian Patent Office or any person approved by the applicant in the individual case.

AUSTRALIA

The applicant hereby gives notice that the furnishing of a sample of a microorganism shall only be effected prior to the grant of a patent, or prior to the lapsing, refusal or withdrawal of the application, to a person who is a skilled addressee without an interest in the invention (Regulation 3.25(3) of the Australian Patents Regulations).

FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

The applicant hereby requests that, until the application has been laid open to public inspection (by the Danish Patent Office), or has been finally decided upon by the Danish Patent Office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Danish Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Danish Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Danish Patent Office or any person by the applicant in the individual case.

SWEDEN

The applicant hereby requests that, until the application has been laid open to public inspection (by the Swedish Patent Office), or has been finally decided upon by the Swedish Patent Office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the International Bureau before the expiration of 16 months from the priority date (preferably on the Form PCT/RO/134 reproduced in annex Z of Volume I of the PCT Applicant's Guide). If such a request has been filed by the applicant any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Swedish Patent Office or any person approved by a applicant in the individual case.

NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

Applicant's or agent's file reference number	PZ031PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>249</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	
Date of deposit June 26, 1998	Accession Number 203027
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States) Europe In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28(4) EPC).	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposits")	
For receiving Office use only <input checked="" type="checkbox"/> This sheet was received with the international application Authorized officer Yusuf E. Sams PCT International Division	For International Bureau use only <input type="checkbox"/> This sheet was received by the International Bureau on: Authorized officer

CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

The applicant hereby requests that the application has been laid open to public inspection (by the Norwegian Patent Office), or has been finally decided upon by the Norwegian Patent Office without having been laid open inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Norwegian Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Norwegian Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on the list of recognized experts drawn up by the Norwegian Patent Office or any person approved by the applicant in the individual case.

AUSTRALIA

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FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

The applicant hereby requests that, until the application has been laid open to public inspection (by the Danish Patent Office), or has been finally decided upon by the Danish Patent office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Danish Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Danish Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Danish Patent Office or any person by the applicant in the individual case.

SWEDEN

The applicant hereby requests that, until the application has been laid open to public inspection (by the Swedish Patent Office), or has been finally decided upon by the Swedish Patent Office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the International Bureau before the expiration of 16 months from the priority date (preferably on the Form PCT/RO/134 reproduced in annex Z of Volume I of the PCT Applicant's Guide). If such a request has been filed by the applicant any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Swedish Patent Office or any person approved by a applicant in the individual case.

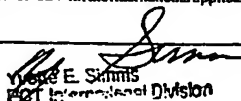
NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

Applicant's or agent's file reference number	PZ031PCT	International application	Unassigned
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>253</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution American Type Culture Collection	
Address of depositary institution (including postal code and country) 10801 University Boulevard Manassas, Virginia 20110-2209 United States of America	
Date of deposit July 27, 1998	Accession Number 203071
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States) Europe In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28(4) EPC).	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable) The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposit")	
<div> <div>For receiving Office use only</div> <div> <input checked="" type="checkbox"/> This sheet was received with the international application </div> <div> Authorized officer  Wayne E. Skimits PCT International Division </div> </div>	<div> <div>For International Bureau use only</div> <div> <input type="checkbox"/> This sheet was received by the International Bureau on: </div> <div> Authorized officer </div> </div>

CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

The applicant hereby requests that the application has been laid open to public inspection (by the Norwegian Patent Office), or has been finally decided upon by the Norwegian Patent Office without having been laid open inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Norwegian Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Norwegian Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on the list of recognized experts drawn up by the Norwegian Patent Office or any person approved by the applicant in the individual case.

AUSTRALIA

The applicant hereby gives notice that the furnishing of a sample of a microorganism shall only be effected prior to the grant of a patent, or prior to the lapsing, refusal or withdrawal of the application, to a person who is a skilled addressee without an interest in the invention (Regulation 3.25(3) of the Australian Patents Regulations).

FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

The applicant hereby requests that, until the application has been laid open to public inspection (by the Danish Patent Office), or has been finally decided upon by the Danish Patent Office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Danish Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Danish Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Danish Patent Office or any person by the applicant in the individual case.

SWEDEN

The applicant hereby requests that, until the application has been laid open to public inspection (by the Swedish Patent Office), or has been finally decided upon by the Swedish Patent Office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the International Bureau before the expiration of 16 months from the priority date (preferably on the Form PCT/RO/134 reproduced in annex Z of Volume I of the PCT Applicant's Guide). If such a request has been filed by the applicant any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Swedish Patent Office or any person approved by a applicant in the individual case.

NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

Applicant's or agent's file reference number	PZ031PCT	International application
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INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13bis)

A. The indications made below relate to the microorganism referred to in the description on page <u>259</u> , line <u>N/A</u>	
B. IDENTIFICATION OF DEPOSIT Further deposits are identified on an additional sheet <input type="checkbox"/>	
Name of depositary institution <u>American Type Culture Collection</u>	
Address of depositary institution (including postal code and country) <u>10801 University Boulevard</u> <u>Manassas, Virginia 20110-2209</u> <u>United States of America</u>	
Date of deposit <u>July 27, 1998</u>	Accession Number <u>203070</u>
C. ADDITIONAL INDICATIONS (leave blank if not applicable) This information is continued on an additional sheet <input type="checkbox"/>	
D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE (if the indications are not for all designated States)	
Europe In respect to those designations in which a European Patent is sought a sample of the deposited microorganism will be made available until the publication of the mention of the grant of the European patent or until the date on which application has been refused or withdrawn or is deemed to be withdrawn, only by the issue of such a sample to an expert nominated by the person requesting the sample (Rule 28(4) EPC).	
E. SEPARATE FURNISHING OF INDICATIONS (leave blank if not applicable)	
The indications listed below will be submitted to the International Bureau later (specify the general nature of the indications e.g., "Accession Number of Deposit")	
<input checked="" type="checkbox"/> For receiving Office use only This sheet was received with the international application Authorized officer <u>Y. E. EMMIS</u> PCT International Division	<input type="checkbox"/> For International Bureau use only This sheet was received by the International Bureau on: Authorized officer

CANADA

The applicant requests that, until either a Canadian patent has been issued on the basis of an application or the application has been refused, or is abandoned and no longer subject to reinstatement, or is withdrawn, the Commissioner of Patents only authorizes the furnishing of a sample of the deposited biological material referred to in the application to an independent expert nominated by the Commissioner, the applicant must, by a written statement, inform the International Bureau accordingly before completion of technical preparations for publication of the international application.

NORWAY

The applicant hereby requests that the application has been laid open to public inspection (by the Norwegian Patent Office), or has been finally decided upon by the Norwegian Patent Office without having been laid open inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Norwegian Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Norwegian Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on the list of recognized experts drawn up by the Norwegian Patent Office or any person approved by the applicant in the individual case.

AUSTRALIA

The applicant hereby gives notice that the furnishing of a sample of a microorganism shall only be effected prior to the grant of a patent, or prior to the lapsing, refusal or withdrawal of the application, to a person who is a skilled addressee without an interest in the invention (Regulation 3.25(3) of the Australian Patents Regulations).

FINLAND

The applicant hereby requests that, until the application has been laid open to public inspection (by the National Board of Patents and Regulations), or has been finally decided upon by the National Board of Patents and Registration without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art.

UNITED KINGDOM

The applicant hereby requests that the furnishing of a sample of a microorganism shall only be made available to an expert. The request to this effect must be filed by the applicant with the International Bureau before the completion of the technical preparations for the international publication of the application.

DENMARK

The applicant hereby requests that, until the application has been laid open to public inspection (by the Danish Patent Office), or has been finally decided upon by the Danish Patent Office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the Danish Patent Office not later than at the time when the application is made available to the public under Sections 22 and 33(3) of the Danish Patents Act. If such a request has been filed by the applicant, any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Danish Patent Office or any person by the applicant in the individual case.

SWEDEN

The applicant hereby requests that, until the application has been laid open to public inspection (by the Swedish Patent Office), or has been finally decided upon by the Swedish Patent Office without having been laid open to public inspection, the furnishing of a sample shall only be effected to an expert in the art. The request to this effect shall be filed by the applicant with the International Bureau before the expiration of 16 months from the priority date (preferably on the Form PCT/RO/134 reproduced in annex Z of Volume I of the PCT Applicant's Guide). If such a request has been filed by the applicant any request made by a third party for the furnishing of a sample shall indicate the expert to be used. That expert may be any person entered on a list of recognized experts drawn up by the Swedish Patent Office or any person approved by a applicant in the individual case.

NETHERLANDS

The applicant hereby requests that until the date of a grant of a Netherlands patent or until the date on which the application is refused or withdrawn or lapsed, the microorganism shall be made available as provided in the 31F(1) of the Patent Rules only by the issue of a sample to an expert. The request to this effect must be furnished by the applicant with the Netherlands Industrial Property Office before the date on which the application is made available to the public under Section 22C or Section 25 of the Patents Act of the Kingdom of the Netherlands, whichever of the two dates occurs earlier.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US99/17130

A. CLASSIFICATION OF SUBJECT MATTER IPC(6) : Please See Extra Sheet. US CL : Please See Extra Sheet. According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) U.S. : 536/23.1, 23.5; 435/69.1, 320.1, 252.3, 325, 6, 7.1; 530/350, 300, 387.1; 514/2 Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) APS, DIALOG - Biotech Files		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	JACOBS, K. A. et al. A Genetic Selection For Isolating cDNAs Encoding Secreted Proteins. Gene. 1997, Vol. 198, pages 289-296, see entire document.	1-23
<input type="checkbox"/> Further documents are listed in the continuation of Box C. <input type="checkbox"/> See patent family annex.		
* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "B" earlier document published on or after the international filing date "L" document which may throw doubt on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "A" document member of the same patent family		
Date of the actual completion of the international search 05 OCTOBER 1999		Date of mailing of the international search report 21 OCT 1999
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230		Authorized officer ELIZABETH C. KEMMERER Telephone No. (703) 308-6196

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US99/17130

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☒ Claims Nos.: 1-23
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

Please See Extra Sheet.

3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US99/17130

A. CLASSIFICATION OF SUBJECT MATTER:
IPC (6):

C12N 1/21, 5/10, 15/11, 15/12, 15/63; A61K 38/16, 38/17; C07K 14/00, 14/435, 16/00; G01N 33/50

A. CLASSIFICATION OF SUBJECT MATTER:
US CL :

536/23.1, 23.5; 435/69.1, 320.1, 252.3, 325, 6, 7.1; 530/350, 300, 387.1; 514/2

BOX I. OBSERVATIONS WHERE CLAIMS WERE FOUND UNSEARCHABLE

2. Where no meaningful search could be carried out, specifically:

All of the claims were unsearchable to the extent that they require reference to sequences from the sequence listing or an ATCC deposit. However, the specific sequence and deposit numbers were replaced in the claims with generic designators X, Y and Z. Therefore, no meaningful search of the sequences or deposits per se can be carried out by this Authority. The subject matter of the claims has been searched only to the extent possible with reference to the balance of the description.